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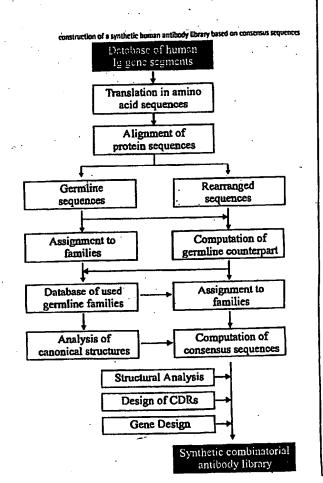
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(54) Title: PROTEIN/(POLY)PEPTIDE LIBRARIES

#### (57) Abstract

The present invention relates to synthetic DNA sequences which encode one or more collections of homologous proteins/(poly)peptides, and methods for generating and applying libraries of these DNA sequences. In particular, the invention relates to the preparation of a library of humanderived antibody genes by the use of synthetic consensus sequences which cover the structural repertoire of antibodies encoded in the human genome. Furthermore, the invention relates to the use of a single consensus antibody gene as a universal framework for highly diverse antibody libraries.



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# Protein/(Poly)peptide Libraries

#### Field of the Invention

The present invention relates to synthetic DNA sequences which encode one or more collections of homologous proteins/(poly)peptides, and methods for generating and applying libraries of these DNA sequences. In particular, the invention relates to the preparation of a library of human-derived antibody genes by the use of synthetic consensus sequences which cover the structural repertoire of antibodies encoded in the human genome. Furthermore, the invention relates to the use of a single consensus antibody gene as a universal framework for highly diverse antibody libraries.

# Background to the Invention

All current recombinant methods which use libraries of proteins/(poly)peptides, e.g. antibodies, to screen for members with desired properties, e.g. binding a given ligand, do not provide the possibility to improve the desired properties of the members in an easy and rapid manner. Usually a library is created either by inserting a random oligonucleotide sequence into one or more DNA sequences cloned from an organism, or a family of DNA sequences is cloned and used as the library. The library is then screened, e.g. using phage display, for members which show the desired property. The sequences of one or more of these resulting molecules are then determined. There is no general procedure available to improve these molecules further on.

Winter (EP 0 368 684 B1) has provided a method for amplifying (by PCR), cloning, and expressing antibody variable region genes. Starting with these genes he was able to create libraries of functional antibody fragments by randomizing the CDR3 of the heavy and/or the light chain. This process is functionally equivalent to the natural process of VJ and VDJ recombination which occurs during the development of B-cells in the immune system.

However the Winter invention does not provide a method for optimizing the binding affinities of antibody fragments further on, a process which would be functionally equivalent to the naturally occurring phenomenon of "affinity maturation", which is provided by the present invention. Furthermore, the Winter invention does not provide for artificial variable region genes, which represent a whole family of

structurally similar natural genes, and which can be assembled from synthetic DNA oligonucleotides. Additionally, Winter does not enable the combinatorial assembly of portions of antibody variable regions, a feature which is provided by the present invention. Furthermore, this approach has the disadvantage that the genes of all antibodies obtained in the screening procedure have to be completely sequenced, since, except for the PCR priming regions, no additional sequence information about the library members is available. This is time and labor intensive and potentially leads to sequencing errors.

The teaching of Winter as well as other approaches have tried to create large antibody libraries having high diversity in the complementarity determining regions (CDRs) as well as in the frameworks to be able to find antibodies against as many different antigens as possible. It has been suggested that a single universal framework may be useful to build antibody libraries, but no approach has yet been successful.

Another problem lies in the production of reagents derived from antibodies. Small antibody fragments show exciting promise for use as therapeutic agents, diagnostic reagents, and for biochemical research. Thus, they are needed in large amounts, and the expression of antibody fragments, e.g. Fv, single-chain Fv (scFv), or Fab in the periplasm of E. coli (Skerra & Plückthun, 1988; Better et al., 1988) is now used routinely in many laboratories. Expression yields vary widely, however. While some fragments yield up to several mg of functional, soluble protein per liter and OD of culture broth in shake flask culture (Carter et al., 1992, Plückthun et al. 1996), other fragments may almost exclusively lead to insoluble material, often found in so-called inclusion bodies. Functional protein may be obtained from the latter in modest yields by a laborious and time-consuming refolding process. The factors influencing antibody expression levels are still only poorly understood. Folding efficiency and stability of the antibody fragments, protease lability and toxicity of the expressed proteins to the host cells often severely limit actual production levels, and several attempts have been tried to increase expression yields. For example, Knappik & Plückthun (1995) could show that expression yield depends on the antibody sequence. They identified key residues in the antibody framework which influence expression yields dramatically. Similarly, Ullrich et al. (1995) found that point mutations in the CDRs can increase the yields in periplasmic antibody fragment expression. Nevertheless, these strategies are only applicable to a few antibodies. Since the Winter invention uses existing repertoires of antibodies, no influence on expressibility of the genes is possible.

Furthermore, the findings of Knappik & Plückthun and Ullrich demonstrate that the knowledge about antibodies, especially about folding and expression is still increasing. The Winter invention does not allow to incorporate such improvements into the library design.

The expressibility of the genes is important for the library quality as well, since the screening procedure relies in most cases on the display of the gene product on a phage surface, and efficient display relies on at least moderate expression of the gene.

These disadvantages of the existing methodologies are overcome by the present invention, which is applicable for all collections of homologous proteins. It has the following novel and useful features illustrated in the following by antibodies as an example:

Artificial antibodies and fragments thereof can be constructed based on known antibody sequences, which reflect the structural properties of a whole group of homologous antibody genes. Therefore it is possible to reduce the number of different genes without any loss in the structural repertoire. This approach leads to a limited set of artificial genes, which can be synthesized de novo, thereby allowing introduction of cleavage sites and removing unwanted cleavages sites. Furthermore, this approach enables (i), adapting the codon usage of the genes to that of highly expressed genes in any desired host cell and (ii), analyzing all possible pairs of antibody light (L) and heavy (H) chains in terms of interaction preference, antigen preference or recombinant expression titer, which is virtually impossible using the complete collection of antibody genes of an organism and all combinations thereof.

The use of a limited set of completely synthetic genes makes it possible to create cleavage sites at the boundaries of encoded structural sub-elements. Therefore, each gene is built up from modules which represent structural sub-elements on the protein/(poly)peptide level. In the case of antibodies, the modules consist of "framework" and "CDR" modules. By creating separate framework and CDR modules, different combinatorial assembly possibilities are enabled. Moreover, if two or more artificial genes carry identical pairs of cleavage sites at the boundaries of each of the genetic sub-elements, pre-built libraries of sub-elements can be inserted in these genes simultaneously, without any additional information related to any particular gene sequence. This strategy enables rapid optimization of, for example, antibody affinity, since DNA cassettes encoding libraries of genetic sub-elements can be (i), pre-built, stored and reused and (ii), inserted in any of these

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sequences at the right position without knowing the actual sequence or having to determine the sequence of the individual library member.

Additionally, new information about amino acid residues important for binding, stability, or solubility and expression could be integrated into the library design by replacing existing modules with modules modified according to the new observations.

The limited number of consensus sequences used for creating the library allows to speed up the identification of binding antibodies after screening. After having identified the underlying consensus gene sequence, which could be done by sequencing or by using fingerprint restriction sites, just those part(s) comprising the random sequence(s) have to be determined. This reduces the probability of sequencing errors and of false-positive results.

The above mentioned cleavage sites can be used only if they are unique in the vector system where the artificial genes have been inserted. As a result, the vector has to be modified to contain none of these cleavage sites. The construction of a vector consisting of basic elements like resistance gene and origin of replication, where cleavage sites have been removed, is of general interest for many cloning attempts. Additionally, these vector(s) could be part of a kit comprising the above mentioned artificial genes and pre-built libraries.

The collection of artificial genes can be used for a rapid humanization procedure of non-human antibodies, preferably of rodent antibodies. First, the amino acid sequence of the non-human, preferably rodent antibody is compared with the amino acid sequences encoded by the collection of artificial genes to determine the most homologous light and heavy framework regions. These genes are then used for insertion of the genetic sub-elements encoding the CDRs of the non-human, preferably rodent antibody.

Surprisingly, it has been found that with a combination of only one consensus sequence for each of the light and heavy chains of a scFv fragment an antibody repertoire could be created yielding antibodies against virtually every antigen. Therefore, one aspect of the present invention is the use of a single consensus sequence as a universal framework for the creation of useful (poly)peptide libraries and antibody consensus sequences useful therefor.

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#### **Detailed Description of the Invention**

The present invention enables the creation of useful libraries of (poly)peptides. In a first embodiment, the invention provides for a method of setting up nucleic acid sequences suitable for the creation of said libraries. In a first step, a collection of at least three homologous proteins is identified and then analyzed. Therefore, a database of the protein sequences is established where the protein sequences are aligned to each other. The database is used to define subgroups of protein sequences which show a high degree of similarity in both the sequence and, if information is available, in the structural arrangement. For each of the subgroups a (poly)peptide sequence comprising at least one consensus sequence is deduced which represents the members of this subgroup; the complete collection of (poly)peptide sequences represent therefore the complete structural repertoire of the collection of homologous proteins. These artificial (poly)peptide sequences are then analyzed, if possible, according to their structural properties to identify unfavorable interactions between amino acids within said (poly)peptide sequences or between said or other (poly)peptide sequences, for example, in multimeric proteins. Such interactions are then removed by changing the consensus sequence accordingly. The (poly)peptide sequences are then analyzed to identify subelements such as domains, loops, helices or CDRs. The amino acid sequence is backtranslated into a corresponding coding nucleic acid sequence which is adapted to the codon usage of the host planned for expressing said nucleic acid sequences. A set of cleavage sites is set up in a way that each of the sub-sequences encoding the sub-elements identified as described above, is flanked by two sites which do not occur a second time within the nucleic acid sequence. This can be achieved by either identifying a cleavage site already flanking a sub-sequence of by changing one or more nucleotides to create the cleavage site, and by removing that site from the remaining part of the gene. The cleavage sites should be common to all corresponding sub-elements or sub-sequences, thus creating a fully modular arrangement of the sub-sequences in the nucleic acid sequence and of the subelements in the corresponding (poly)peptide.

In a further embodiment, the invention provides for a method which sets up two or more sets of (poly)peptides, where for each set the method as described above is performed, and where the cleavage sites are not only unique within each set but also between any two sets. This method can be applied for the creation of (poly)peptide libraries comprising for example two  $\alpha$ -helical domains from two different proteins, where said library is screened for novel hetero-association domains.

In yet a further embodiment, at least two of the sets as described above, are derived from the same collection of proteins or at least a part of it. This describes libraries comprising for example, but not limited to, two domains from antibodies such as VH and VL, or two extracellular loops of transmembrane receptors.

In another embodiment, the nucleic acid sequences set up as described above, are synthesized. This can be achieved by any one of several methods well known to the practitioner skilled in the art, for example, by total gene synthesis or by PCR-based approaches.

In one embodiment, the nucleic acid sequences are cloned into a vector. The vector could be a sequencing vector, an expression vector or a display (e.g. phage display) vector, which are well known to those skilled in the art. Any vector could comprise one nucleic acid sequence, or two or more nucleic sequences, either in different or the same operon. In the last case, they could either be cloned separately or as contiguous sequences.

In one embodiment, the removal of unfavorable interactions as described above, leads to enhanced expression of the modified (poly)peptides.

In a preferred embodiment, one or more sub-sequences of the nucleic acid sequences are replaced by different sequences. This can be achieved by excising the sub-sequences using the conditions suitable for cleaving the cleavage sites adjacent to or at the end of the sub-sequence, for example, by using a restriction enzyme at the corresponding restriction site under the conditions well known to those skilled in the art, and replacing the sub-sequence by a different sequence compatible with the cleaved nucleic acid sequence. In a further preferred embodiment, the different sequences replacing the initial sub-sequence(s) are genomic or rearranged genomic sequences, for example in grafting CDRs from nonhuman antibodies onto consensus antibody sequences for rapid humanization of non-human antibodies. In the most preferred embodiment, the different sequences are random sequences, thus replacing the sub-sequence by a collection of sequences to introduce variability and to create, a library. The random sequences can be assembled in various ways, for example by using a mixture of mononucleotides or preferably a mixture of trinucleotides (Virnekäs et al., 1994) during automated oligonucleotide synthesis, by error-prone PCR or by other methods well known to the practitioner in the art. The random sequences may be completely randomized or biased towards or against certain codons according to

the amino acid distribution at certain positions in known protein sequences. Additionally, the collection of random sub-sequences may comprise different numbers of codons, giving rise to a collection of sub-elements having different lengths.

In another embodiment, the invention provides for the expression of the nucleic acid sequences from a suitable vector and under suitable conditions well known to those skilled in the art.

In a further preferred embodiment, the (poly)peptides expressed from said nucleic acid sequences are screened and, optionally, optimized. Screening may be performed by using one of the methods well known to the practitioner in the art, such as phage-display, selectively infective phage, polysome technology to screen for binding, assay systems for enzymatic activity or protein stability. (Poly)peptides having the desired property can be identified by sequencing of the corresponding nucleic acid sequence or by amino acid sequencing or mass spectrometry. In the case of subsequent optimization, the nucleic acid sequences encoding the initially selected (poly)peptides can optionally be used without sequencing. Optimization is performed by repeating the replacement of sub-sequences by different sequences, preferably by random sequences, and the screening step one or more times.

The desired property the (poly)peptides are screened for is preferably, but not exclusively, selected from the group of optimized affinity or specificity for a target molecule, optimized enzymatic activity, optimized expression yields, optimized stability and optimized solubility.

In one embodiment, the cleavage sites flanking the sub-sequences are sites recognized and cleaved by restriction enzymes, with recognition and cleavage sequences being either identical or different, the restricted sites either having blunt or sticky ends.

The length of the sub-elements is preferably, but not exclusively ranging between 1 amino acid, such as one residue in the active site of an enzyme or a structure-determining residue, and 150 amino acids, as for whole protein domains. Most preferably, the length ranges between 3 and 25 amino acids, such as most commonly found in CDR loops of antibodies.

The nucleic acid sequences could be RNA or, preferably, DNA.

In one embodiment, the (poly)peptides have an amino acid pattern characteristic of a particular species. This can for example be achieved by deducing the consensus sequences from a collection of homologous proteins of just one species, most preferably from a collection of human proteins. Since the (poly)peptides comprising consensus sequences are artificial, they have to be compared to the protein sequence(s) having the closest similarity to ensure the presence of said characteristic amino acid pattern.

In one embodiment, the invention provides for the creation of libraries of (poly)peptides comprising at least part of members or derivatives of the immunoglobulin superfamily, preferably of member or derivatives of the immunoglobulins. Most preferably, the invention provides for the creation of libraries of human antibodies, wherein said (poly)peptides are or are derived from heavy or light chain variable regions wherein said structural sub-elements are framework regions (FR) 1, 2, 3, or 4 or complementary determining regions (CDR) 1, 2, or 3. In a first step, a database of published antibody sequences of human origin is established where the antibody sequences are aligned to each other. The database is used to define subgroups of antibody sequences which show a high degree of similarity in both the sequence and the canonical fold of CDR loops (as determined by analysis of antibody structures). For each of the subgroups a consensus sequence is deduced which represents the members of this subgroup; the complete collection of consensus sequences represent therefore the complete structural repertoire of human antibodies.

These artificial genes are then constructed e.g. by total gene synthesis or by the use of synthetic genetic subunits. These genetic subunits correspond to structural subelements on the (poly)peptide level. On the DNA level, these genetic subunits are defined by cleavage sites at the start and the end of each of the sub-elements, which are unique in the vector system. All genes which are members of the collection of consensus sequences are constructed such that they contain a similar pattern of corresponding genetic sub-sequences. Most preferably, said (poly)peptides are or are derived from the HuCAL consensus genes:  $V_K1$ ,  $V_K2$ ,  $V_K3$ ,  $V_K4$ ,  $V_K1$ ,  $V_K2$ ,  $V_K3$ ,  $V_K4$ ,  $V_K1$ 

This collection of DNA molecules can then be used to create libraries of antibodies or antibody fragments, preferably Fv, disulphide-linked Fv, single-chain Fv (scFv), or Fab fragments, which may be used as sources of specificities against new target antigens. Moreover, the affinity of the antibodies can be optimized using pre-built library cassettes and a general procedure. The invention provides a method for identifying one or more genes encoding one or more antibody fragments which

binds to a target, comprising the steps of expressing the antibody fragments, and then screening them to isolate one or more antibody fragments which bind to a given target molecule. Preferably, an scFv fragment library comprising the combination of HuCAL VH3 and HuCAL V $\lambda$ 2 consensus genes and at least a random sub-sequence encoding the heavy chain CDR3 sub-element is screened for binding antibodies. If necessary, the modular design of the genes can then be used to excise from the genes encoding the antibody fragments one or more genetic sub-sequences encoding structural sub-elements, and replacing them by one or more second sub-sequences encoding structural sub-elements. The expression and screening steps can then be repeated until an antibody having the desired affinity is generated.

Particularly preferred is a method in which one or more of the genetic subunits (e.g. the CDRs) are replaced by a random collection of sequences (the library) using the said cleavage sites. Since these cleavage sites are (i) unique in the vector system and (ii) common to all consensus genes, the same (pre-built) library can be inserted into all artificial antibody genes. The resulting library is then screened against any chosen antigen. Binding antibodies are selected, collected and used as starting material for the next library. Here, one or more of the remaining genetic subunits are randomized as described above.

A further embodiment of the present invention relates to fusion proteins by providing for a DNA sequence which encodes both the (poly)peptide, as described above, as well as an additional moiety. Particularly preferred are moieties which have a useful therapeutic function. For example, the additional moiety may be a toxin molecule which is able to kill cells (Vitetta et al., 1993). There are numerous examples of such toxins, well known to the one skilled in the art, such as the bacterial toxins Pseudomonas exotoxin A, and diphtheria toxin, as well as the plant toxins ricin, abrin, modeccin, saporin, and gelonin. By fusing such a toxin for example to an antibody fragment, the toxin can be targeted to, for example, diseased cells, and thereby have a beneficial therapeutic effect. Alternatively, the additional moiety may be a cytokine, such as IL-2 (Rosenberg & Lotze, 1986), which has a particular effect (in this case a T-cell proliferative effect) on a family of cells. In a further embodiment, the additional moiety may confer on its (poly)peptide partner a means of detection and/or purification. For example, the fusion protein could comprise the modified antibody fragment and an enzyme commonly used for detection purposes, such as alkaline phosphatase (Blake et al., 1984). There are numerous other mojeties which can be used as detection or purification tags, which are well known to the practitioner skilled in the art. Particularly preferred are peptides comprising at least five histidine residues (Hochuli et al., 1988), which are able to bind to metal ions,

and can therefore be used for the purification of the protein to which they are fused (Lindner et al., 1992). Also provided for by the invention are additional moieties such as the commonly used C-myc and FLAG tags (Hopp et al., 1988; Knappik & Plückthun, 1994).

By engineering one or more fused additional domains, antibody fragments or any other (poly)peptide can be assembled into larger molecules which also fall under the scope of the present invention. For example, mini-antibodies (Pack, 1994) are dimers comprising two antibody fragments, each fused to a self-associating dimerization domain. Dimerization domains which are particularly preferred include those derived from a leucine zipper (Pack & Plückthun, 1992) or helix-turn-helix motif (Pack et al., 1993).

All of the above embodiments of the present invention can be effected using standard techniques of molecular biology known to anyone skilled in the art.

In a further embodiment, the random collection of sub-sequences (the library) is inserted into a singular nucleic acid sequence encoding one (poly)peptide, thus creating a (poly)peptide library based on one universal framework. Preferably a random collection of CDR sub-sequences is inserted into a universal antibody framework, for example into the HuCAL  $H3\kappa2$  single-chain Fv fragment described above.

In further embodiments, the invention provides for nucleic acid sequence(s), vector(s) containing the nucleic acid sequence(s), host cell(s) containing the vector(s), and (poly)peptides, obtainable according to the methods described above.

In a further preferred embodiment, the invention provides for modular vector systems being compatible with the modular nucleic acid sequences encoding the (poly)peptides. The modules of the vectors are flanked by restriction sites unique within the vector system and essentially unique with respect to the restriction sites incorporated into the nucleic acid sequences encoding the (poly)peptides, except for example the restriction sites necessary for cloning the nucleic acid sequences into the vector. The list of vector modules comprises origins of single-stranded replication, origins of double-stranded replication for high- and low copy number plasmids, promotor/operator, repressor or terminator elements, resistance genes, potential recombination sites, gene III for display on filamentous phages, signal sequences, purification and detection tags, and sequences of additional moieties.

The vectors are preferably, but not exclusively, expression vectors or vectors suitable for expression and screening of libraries.

In another embodiment, the invention provides for a kit, comprising one or more of the list of nucleic acid sequence(s), recombinant vector(s), (poly)peptide(s), and vector(s) according to the methods described above, and suitable host cell(s) for producing the (poly)peptide(s).

In a preferred embodiment, the invention provides for the creation of libraries of human antibodies. In a first step, a database of published antibody sequences of human origin is established. The database is used to define subgroups of antibody sequences which show a high degree of similarity in both the sequence and the canonical fold (as determined by analysis of antibody structures). For each of the subgroups a consensus sequence is deduced which represents the members of this subgroup; the complete collection of consensus sequences represent therefore the complete structural repertoire of human antibodies.

These artificial genes are then constructed by the use of synthetic genetic subunits. These genetic subunits correspond to structural sub-elements on the protein level. On the DNA level, these genetic subunits are defined by cleavage sites at the start and the end of each of the subelements, which are unique in the vector system. All genes which are members of the collection of consensus sequences are constructed such that they contain a similar pattern of said genetic subunits.

This collection of DNA molecules can then be used to create libraries of antibodies which may be used as sources of specificities against new target antigens. Moreover, the affinity of the antibodies can be optimised using pre-built library cassettes and a general procedure. The invention provides a method for identifying one or more genes encoding one or more antibody fragments which binds to a target, comprising the steps of expressing the antibody fragments, and then screening them to isolate one or more antibody fragments which bind to a given target molecule. If necessary, the modular design of the genes can then be used to excise from the genes encoding the antibody fragments one or more genetic subsequences encoding structural sub-elements, and replacing them by one or more second sub-sequences encoding structural sub-elements. The expression and screening steps can then be repeated until an antibody having the desired affinity is generated.

Particularly preferred is a method in which one or more of the genetic subunits (e.g. the CDR's) are replaced by a random collection of sequences (the library) using the said cleavage sites. Since these cleavage sites are (i) unique in the vector system and (ii) common to all consensus genes, the same (pre-built) library can be inserted into all artificial antibody genes. The resulting library is then screened against any chosen antigen. Binding antibodies are eluted, collected and used as starting material for the next library. Here, one or more of the remaining genetic subunits are randomised as described above.

#### **Definitions**

#### Protein:

The term protein comprises monomeric polypeptide chains as well as homo- or heteromultimeric complexes of two or more polypeptide chains connected either by covalent interactions (such as disulphide bonds) or by non-covalent interactions (such as hydrophobic or electrostatic interactions).

# Analysis of homologous proteins:

The amino acid sequences of three or more proteins are aligned to each other (allowing for introduction of gaps) in a way which maximizes the correspondence between identical or similar amino acid residues at all positions. These aligned sequences are termed homologous if the percentage of the sum of identical and/or similar residues exceeds a defined threshold. This threshold is commonly regarded by those skilled in the art as being exceeded when at least 15% of the amino acids in the aligned genes are identical, and at least 30% are similar. Examples for families of homologous proteins are: immunoglobulin superfamily, scavenger receptor superfamily, fibronectin superfamilies (e.g. type II and III), complement control protein superfamily, cytokine receptor superfamily, cystine knot proteins, tyrosine kinases, and numerous other examples well known to one of ordinary skill in the art.

#### Consensus sequence:

Using a matrix of at least three aligned amino acid sequences, and allowing for gaps in the alignment, it is possible to determine the most frequent amino acid residue at each position. The consensus sequence is that sequence which comprises the amino acids which are most frequently represented at each position. In the event that two or more amino acids are equally represented at a single position, the consensus sequence includes both or all of those amino acids.

### Removing unfavorable interactions:

The consensus sequence is per se in most cases artificial and has to be analyzed in order to change amino acid residues which, for example, would prevent the resulting molecule to adapt a functional tertiary structure or which would block the interaction with other (poly)peptide chains in multimeric complexes. This can be done either by (i) building a three-dimensional model of the consensus sequence using known related structures as a template, and identifying amino acid residues within the model which may interact unfavorably with each other, or (ii) analyzing the matrix of aligned amino acid sequences in order to detect combinations of amino

acid residues within the sequences which frequently occur together in one sequence and are therefore likely to interact with each other. These probable interaction-pairs are then tabulated and the consensus is compared with these "interaction maps". Missing or wrong interactions in the consensus are repaired accordingly by introducing appropriate changes in amino acids which minimize unfavorable interactions.

#### Identification of structural sub-elements:

Structural sub-elements are stretches of amino acid residues within a protein/(poly)peptide which correspond to a defined structural or functional part of the molecule. These can be loops (e.g. CDR loops of an antibody) or any other secondary or functional structure within the protein/(poly)peptide (domains,  $\alpha$ -helices,  $\beta$ -sheets, framework regions of antibodies, etc.). A structural sub-element can be identified using known structures of similar or homologous (poly)peptides, or by using the above mentioned matrices of aligned amino acid sequences. Here the variability at each position is the basis for determining stretches of amino acid residues which belong to a structural sub-element (e.g. hypervariable regions of an antibody).

#### Sub-sequence:

A sub-sequence is defined as a genetic module which is flanked by unique cleavage sites and encodes at least one structural sub-element. It is not necessarily identical to a structural sub-element.

#### Cleavage site:

A short DNA sequence which is used as a specific target for a reagent which cleaves DNA in a sequence-specific manner (e.g. restriction endonucleases).

#### Compatible cleavage sites:

Cleavage sites are compatible with each other, if they can be efficiently ligated without modification and, preferably, also without adding an adapter molecule..

# Unique cleavage sites:

A cleavage site is defined as unique if it occurs only once in a vector containing at least one of the genes of interest, or if a vector containing at least one of the genes of interest could be treated in a way that only one of the cleavage sites could be used by the cleaving agent.

# Corresponding (poly)peptide sequences:

Sequences deduced from the same part of one group of homologous proteins are called corresponding (poly)peptide sequences.

#### Common cleavage sites:

A cleavage site in at least two corresponding sequences, which occurs at the same functional position (i.e. which flanks a defined sub-sequence), which can be hydrolyzed by the same cleavage tool and which yields identical compatible ends is termed a common cleavage site.

# Excising genetic sub-sequences:

A method which uses the unique cleavage sites and the corresponding cleavage reagents to cleave the target DNA at the specified positions in order to isolate, remove or replace the genetic sub-sequence flanked by these unique cleavage sites.

# Exchanging genetic sub-sequences:

A method by which an existing sub-sequence is removed using the flanking cleavage sites of this sub-sequence, and a new sub-sequence or a collection of sub-sequences, which contain ends compatible with the cleavage sites thus created, is inserted.

#### **Expression of genes:**

The term expression refers to in vivo or in vitro processes, by which the information of a gene is transcribed into mRNA and then translated into a protein/(poly)peptide. Thus, the term expression refers to a process which occurs inside cells, by which the information of a gene is transcribed into mRNA and then into a protein. The term expression also includes all events of post-translational modification and transport, which are necessary for the (poly)peptide to be functional.

#### Screening of protein/(poly)peptide libraries:

Any method which allows isolation of one or more proteins/(poly)peptides having a desired property from other proteins/(poly)peptides within a library.

# Amino acid pattern characteristic for a species:

A (poly)peptide sequence is assumed to exhibit an amino acid pattern characteristic for a species if it is deduced from a collection of homologous proteins from just this species.

#### Immunoglobulin superfamily (IgSF):

The IgSF is a family of proteins comprising domains being characterized by the immunoglobulin fold. The IgSF comprises for example T-cell receptors and the immunoglobulins (antibodies).

# Antibody framework:

A framework of an antibody variable domain is defined by Kabat et al. (1991) as the part of the variable domain which serves as a scaffold for the antigen binding loops of this variable domain.

### **Antibody CDR:**

The CDRs (complementarity determining regions) of an antibody consist of the antigen binding loops, as defined by Kabat et al. (1991). Each of the two variable domains of an antibody Fv fragment contain three CDRs.

#### **HuCAL**:

Acronym for <u>Human Combinatorial Antibody Library</u>. Antibody Library based on modular consensus genes according to the invention (see Example 1).

#### Antibody fragment:

Any portion of an antibody which has a particular function, e.g. binding of antigen. Usually, antibody fragments are smaller than whole antibodies. Examples are Fv, disulphide-linked Fv, single-chain Fv (scFv), or Fab fragments. Additionally, antibody fragments are often engineered to include new functions or properties.

#### Universal framework:

One single framework which can be used to create the full variability of functions, specificities or properties which is originally sustained by a large collection of different frameworks, is called universal framework.

# Binding of an antibody to its target:

The process which leads to a tight and specific association between an antibody and a corresponding molecule or ligand is called binding. A molecule or ligand or any part of a molecule or ligand which is recognized by an antibody is called the target.

# Replacing genetic sub-sequences

A method by which an existing sub-sequence is removed using the flanking cleavage sites of this sub-sequence, and a new sub-sequence or collection of sub-

sequences, which contains ends compatible with the cleavage sites thus created, is inserted.

# Assembling of genetic sequences:

Any process which is used to combine synthetic or natural genetic sequences in a specific manner in order to get longer genetic sequences which contain at least parts of the used synthetic or natural genetic sequences.

# Analysis of homologous genes:

The corresponding amino acid sequences of two or more genes are aligned to each other in a way which maximizes the correspondence between identical or similar amino acid residues at all positions. These aligned sequences are termed homologous if the percentage of the sum of identical and/or similar residues exceeds a defined threshold. This threshold is commonly regarded by those skilled in the art as being exceeded when at least 15 per cent of the amino acids in the aligned genes are identical, and at least 30 per cent are similar.

# Legends to Figures and Tables

Fig. 1: Flow chart outlining the process of construction of a synthetic human antibody library based on consensus sequences.

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- Fig. 10: Sequencing results of independent clones from the initial library, translated into the corresponding amino acid sequences. (A) Amino acid sequence of the VH3 consensus heavy chain CDR3 (position 93 to 102, Kabat numbering). (B) Amino acid sequences of 12 clones of the 10-mer library. (C) Amino acid sequences of 11 clones of the 15-mer library, \*: single base deletion.
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- Fig. 25: Schematic representation of the modular pCAL vector system.
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Fig. 27: Functional map and sequence of the multi-cloning site module (MCS)

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- Fig. 40: Expression of all 49 HuCAL scFvs obtained by combining each of the 7 VH genes with each of the 7 VL genes (pBS13, 30°C): Values are given for the percentage of soluble vs. insoluble material, the total and the soluble amount compared to the combination H3κ2, which was set to 100%. In addition, the corresponding values for the McPC603 scFv are given.
- Table 1: Summary of human immunoglobulin germline sequences used for computing the germline membership of rearranged sequences. (A) kappa sequences, (B) lambda sequences and (C), heavy chain sequences. (1) The germline name used in the various calculations, (2) the references number for the corresponding sequence (see appendix for sequence related citations), (3) the family where each sequence belongs to and (4), the various names found in literature for germline genes with identical amino acid sequences.
- Table 2: Rearranged human sequences used for the calculation of consensus sequences. (A) kappa sequences, (B) lambda sequences and (C), heavy chain sequences. The table summarized the name of the sequence (1),

the length of the sequence in amino acids (2), the germline family (3) as well as the computed germline counterpart (4). The number of amino acid exchanges between the rearranged sequence and the germline sequence is tabulated in (5), and the percentage of different amino acids is given in (6). Column (7) gives the references number for the corresponding sequence (see appendix for sequence related citations).

- Table 3: Assignment of rearranged V sequences to their germline counterparts.

  (A) kappa sequences, (B) lambda sequences and (C), heavy chain sequences. The germline genes are tabulated according to their family (1), and the number of rearranged genes found for every germline gene is given in (2).
- Table 4: Computation of the consensus sequence of the rearranged V kappa sequences. (A), V kappa subgroup 1, (B), V kappa subgroup 2, (C), V kappa subgroup 3 and (D), V kappa subgroup 4. The number of each amino acid found at each position is tabulated together with the statistical analysis of the data. (1) Amino acids are given with their standard one-letter abbreviations (and B means D or N, Z means E or Q and X means any amino acid). The statistical analysis summarizes the number of sequences found at each position (2), the number of occurrences of the most common amino acid (3), the amino acid residue which is most common at this position (4), the relative frequency of the occurrence of the most common amino acid (5) and the number of different amino acids found at each position (6).
- Table 5: Computation of the consensus sequence of the rearranged V lambda sequences. (A), V lambda subgroup 1, (B), V lambda subgroup 2, and (C), V lambda subgroup 3. The number of each amino acid found at each position is tabulated together with the statistical analysis of the data. Abbreviations are the same as in Table 4.
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#### Examples

# Example 1: Design of a Synthetic Human Combinatorial Antibody Library (HuCAL)

The following example describes the design of a fully synthetic human combinatorial antibody library (HuCAL), based on consensus sequences of the human immunoglobulin repertoire, and the synthesis of the consensus genes. The general procedure is outlined in Fig. 1.

# 1.1 Sequence database

# 1.1.1 Collection and alignment of human immunoglobulin sequences

In a first step, sequences of variable domains of human immunoglobulins have been collected and divided into three sub bases: V heavy chain (VH), V kappa (V $\kappa$ ) and V lambda (V $\lambda$ ). For each sequence, the gene sequence was then translated into the corresponding amino acid sequence. Subsequently, all amino acid sequences were aligned according to Kabat et al. (1991). In the case of V $\lambda$  sequences, the numbering system of Chuchana et al. (1990) was used. Each of the three main databases was then divided into two further sub bases: the first sub base contained all sequences derived from rearranged V genes, where more than 70 positions of the sequence were known. The second sub base contained all germline gene segments (without the D- and J- minigenes; pseudogenes with internal stop codons were also removed). In all cases, where germline sequences with identical amino acid sequence but different names were found, only one sequence was used (see Table 1). The final databases of rearranged sequences contained 386, 149 and 674 entries for V $\kappa$ , V $\lambda$  and VH, respectively. The final databases of germline sequences contained 48, 26 and 141 entries for V $\kappa$ , V $\lambda$  and VH, respectively.

# 1.1.2 Assignment of sequences to subgroups

The sequences in the three germline databases where then grouped according to sequence homology (see also Tomlinson et al., 1992, Williams & Winter, 1993, and Cox et al., 1994). In the case of  $V\kappa$ , 7 families could be established.  $V\lambda$  was divided into 8 families and VH into 6 families. The VH germline genes of the VH7 family (Van Dijk et al., 1993) were grouped into the VH1 family, since the genes of the two families are highly homologous. Each family contained different numbers of germline genes, varying from 1 (for example VH6) to 47 (VH3).

# 1.2 Analysis of sequences

# 1.2.1 Computation of germline membership

For each of the 1209 amino acid sequences in the databases of rearranged genes, the nearest germline counterpart, i.e. the germline sequence with the smallest number of amino acid differences was then calculated. After the germline counterpart was found, the number of somatic mutations which occurred in the rearranged gene and which led to amino acid exchanges could be tabulated. In 140 cases, the germline counterpart could not be calculated exactly, because more than one germline gene was found with an identical number of amino acid exchanges. These rearranged sequences were removed from the database. In a few cases, the number of amino acid exchanges was found to be unusually large (>20 for VL and >25 for VH), indicating either heavily mutated rearranged genes or derivation from germline genes not present in the database. Since it was not possible to distinguish between these two possibilities, these sequences were also removed from the database. Finally, 12 rearranged sequences were removed from the database because they were found to have very unusual CDR lengths and composition or unusual amino acids at canonical positions (see below). In summary, 1023 rearranged sequences out of 1209 (85%) could be clearly assigned to their germline counterparts (see Table 2).

After this calculation, every rearranged gene could be arranged in one of the families established for the germline genes. Now the usage of each germline gene, i.e. the number of rearranged genes which originate from each germline gene, could be calculated (see Table 2). It was found that the usage was strongly biased towards a subset of germline genes, whereas most of the germline genes were not present as rearranged genes in the database and therefore apparently not used in the immune system (Table 3). This observation had already been reported in the case of  $V\kappa$  (Cox, et al., 1994). All germline gene families, where no or only very few rearranged counterparts could be assigned, were removed from the database, leaving 4  $V\kappa$ , 3  $V\lambda$ , and 6 VH families.

# 1.2.2 Analysis of CDR conformations

The conformation of the antigen binding loops of antibody molecules, the CDRs, is strongly dependent on both the length of the CDRs and the amino acid residues located at the so-called canonical positions (Chothia & Lesk, 1987). It has been found that only a few canonical structures exist, which determine the structural

repertoire of the immunoglobulin variable domains (Chothia et al., 1989). The canonical amino acid positions can be found in CDR as well as framework regions. The 13 used germline families defined above (7 VL and 6 VH) were now analyzed for their canonical structures in order to define the structural repertoire encoded in these families.

In 3 of the 4 V $\kappa$  families (V $\kappa$ 1, 2 and 4), one different type of CDR1 conformation could be defined for every family. The family V $\kappa$ 3 showed two types of CDR1 conformation: one type which was identical to V $\kappa$ 1 and one type only found in V $\kappa$ 3. All V $\kappa$  CDR2s used the same type of canonical structure. The CDR3 conformation is not encoded in the germline gene segments. Therefore, the 4 V $\kappa$  families defined by sequence homology and usage corresponded also to 4 types of canonical structures found in V $\kappa$  germline genes.

The 3 V $\lambda$  families defined above showed 3 types of CDR1 conformation, each family with one unique type. The V $\lambda$ 1 family contained 2 different CDR1 lengths (13 and 14 amino acids), but identical canonical residues, and it is thought that both lengths adopt the same canonical conformation (Chothia & Lesk, 1987). In the CDR2 of the used V $\lambda$  germlines, only one canonical conformation exists, and the CDR3 conformation is not encoded in the germline gene segments. Therefore, the 3 V $\lambda$ 4 families defined by sequence homology and usage corresponded also to 3 types of canonical structures.

The structural repertoire of the human VH sequences was analyzed in detail by Chothia et al., 1992. In total, 3 conformations of CDR1 (H1-1, H1-2 and H1-3) and 6 conformations of CDR2 (H2-1, H2-2, H2-3, H2-4, H2-5 and H2-x) could be defined. Since the CDR3 is encoded in the D- and J-minigene segments, no particular canonical residues are defined for this CDR.

All the members of the VH1 family defined above contained the CDR1 conformation H1-1, but differed in their CDR2 conformation: the H2-2 conformation was found in 6 germline genes, whereas the conformation H2-3 was found in 8 germline genes. Since the two types of CDR2 conformations are defined by different types of amino acid at the framework position 72, the VH1 family was divided into two subfamilies: VH1A with CDR2 conformation H2-2 and VH1B with the conformation H2-3. The members of the VH2 family all had the conformations H1-3 and H2-1 in CDR1 and CDR2, respectively. The CDR1 conformation of the VH3 members was found in all cases to be H1-1, but 4 different types were found in CDR2 (H2-1, H2-3, H2-4 and H2-x). In these CDR2 conformations, the canonical framework residue 71 is always

defined by an arginine. Therefore, it was not necessary to divide the VH3 family into subfamilies, since the 4 types of CDR2 conformations were defined solely by the CDR2 itself. The same was true for the VH4 family. Here, all 3 types of CDR1 conformations were found, but since the CDR1 conformation was defined by the CDR itself (the canonical framework residue 26 was found to be glycine in all cases), no subdivisions were necessary. The CDR2 conformation of the VH4 members was found to be H2-1 in all cases. All members of the VH5 family were found to have the conformation H1-1 and H2-2, respectively. The single germline gene of the VH6 family had the conformations H1-3 and H2-5 in CDR1 and CDR2, respectively.

In summary, all possible CDR conformations of the  $V\kappa$  and  $V\lambda$  genes were present in the 7 families defined by sequence comparison. From the 12 different CDR conformations found in the used VH germline genes, 7 could be covered by dividing the family VH1 into two subfamilies, thereby creating 7 VH families. The remaining 5 CDR conformations (3 in the VH3 and 2 in the VH4 family) were defined by the CDRs themselves and could be created during the construction of CDR libraries. Therefore, the structural repertoire of the used human V genes could be covered by 49 (7 x 7) different frameworks.

# 1.2.3 Computation of consensus sequences

The 14 databases of rearranged sequences (4 Vκ, 3 Vλ and 7 VH) were used to compute the HuCAL consensus sequences of each subgroup (4 HuCAL- Vk. 3 HuCAL- Vλ, 7 HuCAL- VH, see Table 4, 5 and 6). This was done by counting the number of amino acid residues used at each position (position variability) and subsequently identifying the amino acid residue most frequently used at each position. By using the rearranged sequences instead of the used germline sequences for the calculation of the consensus, the consensus was weighted according to the frequency of usage. Additionally, frequently mutated and highly conserved positions could be identified. The consensus sequences were crosschecked with the consensus of the germline families to see whether the rearranged sequences were biased at certain positions towards amino acid residues which do not occur in the collected germline sequences, but this was found not to be the case. Subsequently, the number of differences of each of the 14 consensus sequences to each of the germline sequences found in each specific family was calculated. The overall deviation from the most homologous germline sequence was found to be 2.4 amino acid residues (s.d. = 2.7), ensuring that the "artificial" consensus sequences

can still be considered as truly human sequences as far as immunogenicity is concerned.

#### 1.3 Structural analysis

So far, only sequence information was used to design the consensus sequences. Since it was possible that during the calculation certain artificial combinations of amino acid residues have been created, which are located far away in the sequence but have contacts to each other in the three dimensional structure, leading to destabilized or even misfolded frameworks, the 14 consensus sequences were analyzed according to their structural properties.

It was rationalized that all rearranged sequences present in the database correspond to functional and therefore correctly folded antibody molecules. Hence, the most homologous rearranged sequence was calculated for each consensus sequence. The positions where the consensus differed from the rearranged sequence were identified as potential "artificial residues" and inspected.

The inspection itself was done in two directions. First, the local sequence stretch around each potentially "artificial residue" was compared with the corresponding stretch of all the rearranged sequences. If this stretch was found to be truly artificial, i.e. never occurred in any of the rearranged sequences, the critical residue was converted into the second most common amino acid found at this position and analyzed again. Second, the potentially "artificial residues" were analyzed for their long range interactions. This was done by collecting all available structures of human antibody variable domains from the corresponding PDB files and calculating for every structure the number and type of interactions each amino acid residue established to each side-chain. These "interaction maps" were used to analyze the probable side-chain/side-chain interactions of the potentially "artificial residues". As a result of this analysis, the following residues were exchanged (given is the name of the gene, the position according to Kabat's numbering scheme, the amino acid found at this position as the most abundant one and the amino acid which was used instead):

VH2: S<sub>65</sub>T

Vκ1: N<sub>34</sub>A,

Vκ3: G<sub>9</sub>A, D<sub>60</sub>A, R<sub>77</sub>S

Vλ3: V<sub>78</sub>T

# 1.4 Design of CDR sequences

The process described above provided the complete consensus sequences derived solely from the databases of rearranged sequences. It was rationalized that the CDR1 and CDR2 regions should be taken from the databases of used germline sequences, since the CDRs of rearranged and mutated sequences are biased towards their particular antigens. Moreover, the germline CDR sequences are known to allow binding to a variety of antigens in the primary immune response, where only CDR3 is varied. Therefore, the consensus CDRs obtained from the calculations described above were replaced by germline CDRs in the case of VH and  $V_K$ . In the case of  $V_K$ , a few amino acid exchanges were introduced in some of the chosen germline CDRs in order to avoid possible protease cleavage sites as well as possible structural constraints.

The CDRs of following germline genes have been chosen:

HuCAL gene	CDR1	CDR2
HuCAL-VH1A	VH1-12-1	VH1-12-1
HuCAL-VH1B	VH1-13-16	VH1-13-6,-7,-8,-9
HuCAL-VH2	VH2-31-10,-11,-12,-13	VH2-31-3,-4
HuCAL-VH3	VH3-13-8,-9,-10	VH3-13-8,-9,-10
HuCAL-VH4	VH4-11-7 to -14	VH4-11-8,-9,-11,-12,-14,-16
		VH4-31-17,-18,-19,-20
HuCAL-VH5	VH5-12-1,-2	VH5-12-1,-2
HuCAL-VH6	VH6-35-1	VH6-35-1
HuCAL-Vκ1	Vκ1-14,-15	Vĸ1-2,-3,-4,-5,-7,-8,-12,-13,-18,-19
HuCAL-Vκ2	Vĸ2-6	Vκ2-6
HuCAL-Vk3	Vκ3-1,-4	Vκ3-4
HuCAL-V <sub>K</sub> 4	Vĸ4-1	Vκ4-1
HuCAL-Vλ1	HUMLV117,DPL5	DPL5
HuCAL-Vλ2	DPL11,DPL12	DPL12
HuCAL-Vλ3	DPL23	HUMLV318

In the case of the CDR3s, any sequence could be chosen since these CDRs were planned to be the first to be replaced by oligonucleotide libraries. In order to study the expression and folding behavior of the consensus sequences in *E. coli*, it would be useful to have all sequences with the same CDR3, since the influence of the CDR3s on the folding behavior would then be identical in all cases. The dummy sequences QQHYTTPP and ARWGGDGFYAMDY were selected for the VL chains (kappa and lambda) and for the VH chains, respectively. These sequences are known to be compatible with antibody folding in *E. coli* (Carter et al., 1992).

# 1.5 Gene design

The final outcome of the process described above was a collection of 14 HuCAL amino acid sequences, which represent the frequently used structural antibody repertoire of the human immune system (see Figure 2). These sequences were back-translated into DNA sequences. In a first step, the back-translation was done using only codons which are known to be frequently used in E. coli. These gene sequences were then used for creating a database of all possible restriction endonuclease sites, which could be introduced without changing the corresponding amino acid sequences. Using this database, cleavage sites were selected which were located at the flanking regions of all sub-elements of the genes (CDRs and framework regions) and which could be introduced in all HuCAL VH, V $\kappa$  or V $\lambda$ genes simultaneously at the same position. In a few cases it was not possible to find cleavage sites for all genes of a subgroup. When this happened, the amino acid sequence was changed, if this was possible according to the available sequence and structural information. This exchange was then analyzed again as described above. In total, the following 6 amino acid residues were exchanged during this design (given is the name of the gene, the position according to Kabat's numbering scheme, the amino acid found at this position as the most abundant one and the amino acid which was used instead):

VH2: T<sub>3</sub>Q

VH6: S<sub>42</sub>G

Vκ3: E,D, I<sub>58</sub>V

Vκ4: Κ<sub>24</sub>R

Vλ3: T<sub>22</sub>S

In one case (5'-end of VH framework 3) it was not possible to identify a single cleavage site for all 7 VH genes. Two different type of cleavage sites were used instead: BstEll for HuCAL VH1A, VH1B, VH4 and VH5, and NspV for HuCAL VH2, VH3, VH4 and VH6.

Several restriction endonuclease sites were identified, which were not located at the flanking regions of the sub-elements but which could be introduced in every gene of a given group without changing the amino acid sequence. These cleavage sites were also introduced in order to make the system more flexible for further improvements. Finally, all but one remaining restriction endonuclease sites were removed in every gene sequence. The single cleavage site, which was not removed was different in all genes of a subgroup and could be therefore used as a "fingerprint" site to ease the identification of the different genes by restriction digest. The designed genes, together with the corresponding amino acid sequences and the group-specific restriction endonuclease sites are shown in Figure 3, 4 and 5, respectively.

# 1.6 Gene synthesis and cloning

The consensus genes were synthesized using the method described by Prodromou & Pearl, 1992, using the oligonucleotides shown in Fig. 6. Gene segments encoding the human constant domains  $C\kappa$ ,  $C\lambda$  and CH1 were also synthesized, based on sequence information given by Kabat et al., 1991 (see Fig. 6 and Fig. 7). Since for both the CDR3 and the framework 4 gene segments identical sequences were chosen in all HuCAL  $V\kappa$ ,  $V\lambda$  and VH genes, respectively, this part was constructed only once, together with the corresponding gene segments encoding the constant domains. The PCR products were cloned into pCR-Script KS(+) (Stratagene, Inc.) or pZErO-1 (Invitrogen, Inc.) and verified by sequencing.

# Example 2: Cloning and Testing of a HuCAL-Based Antibody Library

A combination of two of the synthetic consensus genes was chosen after construction to test whether binding antibody fragments can be isolated from a library based on these two consensus frameworks. The two genes were cloned as a single-chain Fv (scFv) fragment, and a VH-CDR3 library was inserted. In order to test the library for the presence of functional antibody molecules, a selection procedure

was carried out using the small hapten fluorescein bound to BSA (FITC-BSA) as antigen.

# 2.1 Cloning of the HuCAL VH3-Vk2 scFv fragment

In order to test the design of the consensus genes, one randomly chosen combination of synthetic light and heavy gene (HuCAL-Vk2 and HuCAL-VH3) was used for the construction of a single-chain antibody (scFv) fragment. Briefly, the gene segments encoding the VH3 consensus gene and the CH1 gene segment including the CDR3 - framework 4 region, as well as the Vκ2 consensus gene and the Cκ gene segment including the CDR3 - framework 4 region were assembled yielding the gene for the VH3-CH1 Fd fragment and the gene encoding the Vκ2-Cκ light chain, respectively. The CH1 gene segment was then replaced by an oligonucleotide cassette encoding a 20-mer peptide linker with the sequence AGGGSGGGGGGGGGGG. The two oligonucleotides encoding this linker were 5'- TCAGCGGGTGGCGGTTCTGGCGGCGGTGGAGCGGTGGCGGTGGTTC-TGGCGGTGGTGCTTCCGATATCGGTCCACGTACGG-3' and 5'-AATTCCGTACG-TGGACCGATATCGGAACCACCACCGCCAGAACCACCGCCACCGCTCCCACCGC CGCCAGAACCGCCACCGC-3', respectively. Finally, the HuCAL-Vk2 gene was inserted via EcoRV and BsiWI into the plasmid encoding the HuCAL-VH3-linker fusion, leading to the final gene HuCAL-VH3-Vk2, which encoded the two consensus sequences in the single-chain format VH-linker-VL. The complete coding sequence is shown in Fig. 8.

# 2.2 Construction of a monovalent phage-display phagemid vector pIG10.3

Phagemid plG10.3 (Fig. 9) was constructed in order to create a phage-display system (Winter et al., 1994) for the H3κ2 scFv gene. Briefly, the EcoRI/HindIII restriction fragment in the phagemid vector plG10 (Ge et al., 1995) was replaced by the c-myc followed by an amber codon (which encodes an glutamate in the amber-suppresser strain XL1 Blue and a stop codon in the non-suppresser strain JM83) and a truncated version of the gene III (fusion junction at codon 249, see Lowman et al., 1991) through PCR mutagenesis.

### 2.3 Construction of H-CDR3 libraries

Heavy chain CDR3 libraries of two lengths (10 and 15 amino acids) were constructed using trinucleotide codon containing oligonucleotides (Virnekäs et al., 1994) as templates and the oligonucleotides complementing the flanking regions as primers. To concentrate only on the CDR3 structures that appear most often in functional antibodies, we kept the salt-bridge of  $R_{\rm H94}$  and  $D_{\rm H101}$  in the CDR3 loop. For the 15-mer library, both phenylalanine and methionine were introduced at position 100 since these two residues were found to occur quite often in human CDR3s of this length (not shown). For the same reason, valine and tyrosine were introduced at position 102. All other randomized positions contained codons for all amino acids except cystein, which was not used in the trinucleotide mixture.

The CDR3 libraries of lengths 10 and 15 were generated from the PCR fragments using oligonucleotide templates O3HCDR103T (5'- GATACGGCCGTGTATTA-TTGCGCGCGT (TRI)6GATTATTGGGGCCAAGGCACCCTG-3') and O3HCDR153T (5'-GATACGGCCGT GTATTATTGCGCGCGT(TRI)10(TTT/ATG)GAT(GTT/TAT)TGGG-GCCAAGGCACCCTG-3'), and primers O3HCDR35 (5'-GATACGGCCGTGTATTA-TTGC-3') and O3HCDR33 (5'-CAGGGTGCCTTGGCCCC-3'), where TRI are trinucleotide mixtures representing all amino acids without cystein, (TTT/ATG) and (GTT/TAT) are trinucleotide mixtures the encodina amino acids phenylalanine/methionine and valine/tyrosine, respectively. The potential diversity of these libraries was  $4.7 \times 10^7$  and  $3.4 \times 10^{10}$  for 10-mer and 15-mer library, respectively. The library cassettes were first synthesized from PCR amplification of the oligo templates in the presence of both primers: 25 pmol of the oligo template O3HCDR103T or O3HCDR153T, 50 pmol each of the primers O3HCDR35 and O3HCDR33, 20 nmol of dNTP, 10x buffer and 2.5 units of Pfu DNA polymerase (Stratagene) in a total volume of 100 µl for 30 cycles (1 minute at 92°C, 1 minute at 62°C and 1 minute at 72°C). A hot-start procedure was used. The resulting mixtures were phenol-extracted, ethanol-precipitated and digested overnight with Eagl and Styl. The vector plG10.3-scH3x2cat, where the Eagl-Styl fragment in the vector pIG10.3-scH3κ2 encoding the H-CDR3 was replaced by the chloramphenicol acetyltransferase gene (cat) flanked with these two sites, was similarly digested. The digested vector (35  $\mu$ g) was gel-purified and ligated with 100  $\mu$ g of the library cassette overnight at 16°C. The ligation mixtures were isopropanol precipitated, airdried and the pellets were redissolved in 100  $\mu l$  of ddH2O. The ligation was mixed with 1 ml of freshly prepared electrocompetent XL1 Blue on ice. 20 rounds of electroporation were performed and the transformants were diluted in SOC medium, shaken at 37°C for 30 minutes and plated out on large LB plates (Amp/Tet/Glucose)

at 37°C for 6-9 hrs. The number of transformants (library size) was 3.2x10<sup>7</sup> and 2.3x10<sup>7</sup> for the 10-mer and the 15-mer library, respectively. The colonies were suspended in 2xYT medium (Amp/Tet/Glucose) and stored as glycerol culture.

In order to test the quality of the initial library, phagemids from 24 independent colonies (12 from the 10-mer and 12 from the 15-mer library, respectively) were isolated and analyzed by restriction digestion and sequencing. The restriction analysis of the 24 phagemids indicated the presence of intact vector in all cases. Sequence analysis of these clones (see Fig. 10) indicated that 22 out of 24 contained a functional sequence in their heavy chain CDR3 regions. 1 out of 12 clones of the 10-mer library had a CDR3 of length 9 instead of 10, and 2 out of 12 clones of the 15-mer library had no open reading frame, thereby leading to a nonfunctional scFv; one of these two clones contained two consecutive inserts, but out of frame (data not shown). All codons introduced were presented in an even distribution.

Expression levels of individual library members were also measured. Briefly, 9 clones from each library were grown in 2xYT medium containing Amp/Tet/0.5% glucose at 37°C overnight. Next day, the cultures were diluted into fresh medium with Amp/Tet. At an OD<sub>600nm</sub> of 0.4, the cultures were induced with 1 mM of IPTG and shaken at RT overnight. Then the cell pellets were suspended in 1 ml of PBS buffer + 1 mM of EDTA. The suspensions were sonicated and the supernatants were separated on an SDS-PAGE under reducing conditions, blotted on nylon membrane and detected with anti-FLAG M1 antibody (see Fig. 11). From the nine clones of the 10-mer library, all express the scFv fragments. Moreover, the gene III / scFv fusion proteins were present in all cases. Among the nine clones from the 15-mer library analyzed, 6/9 (67%) led to the expression of both scFv and the gene III/scFv fusion proteins. More importantly, all clones expressing the scFvs and gene III/scFv fusions gave rise to about the same level of expression.

# 2.4 Biopanning

Phages displaying the antibody libraries were prepared using standard protocols. Phages derived from the 10-mer library were mixed with phages from the 15-mer library in a ratio of 20:1 ( $1x10^{10}$  cfu/well of the 10-mer and  $5x10^8$  cfu/well of the 15-mer phages, respectively). Subsequently, the phage solution was used for panning in ELISA plates (Maxisorp, Nunc) coated with FITC-BSA (Sigma) at concentration of  $100 \ \mu g/ml$  in PBS at 4°C overnight. The antigen-coated wells were blocked with 3% powder milk in PBS and the phage solutions in 1% powder milk were added to each

well and the plate was shaken at RT for 1 hr. The wells were then washed with PBST and PBS (4 times each with shaking at RT for 5 minutes). The bound phages were eluted with 0.1 M triethylamine (TEA) at RT for 10 minutes. The eluted phage solutions were immediately neutralized with 1/2 the volume of 1 M Tris-Cl, pH 7.6. Eluted phage solutions (ca. 450  $\mu$ l) were used to infect 5 ml of XL1 Blue cells at 37°C for 30 min. The infected cultures were then plated out on large LB plates (Amp/Tet/Glucose) and allowed to grow at 37°C until the colonies were visible. The colonies were suspended in 2xYT medium and the glycerol cultures were made as above described. This panning round was repeated twice, and in the third round elution was carried out with addition of fluorescein in a concentration of 100  $\mu$ g/ml in PBS. The enrichment of specific phage antibodies was monitored by panning the initial as well as the subsequent fluorescein-specific sub-libraries against the blocking buffer (Fig. 12). Antibodies with specificity against fluorescein were isolated after 3 rounds of panning.

# 2.5 ELISA measurements

One of the criteria for the successful biopanning is the isolation of individual phage clones that bind to the targeted antigen or hapten. We undertook the isolation of anti-FITC phage antibody clones and characterized them first in a phage ELISA format. After the 3rd round of biopanning (see above), 24 phagemid containing clones were used to inoculate 100  $\mu$ l of 2xYT medium (Amp/Tet/Glucose) in an ELISA plate (Nunc), which was subsequently shaken at 37°C for 5 hrs. 100  $\mu$ l of 2xYT medium (Amp/Tet/1 mM IPTG) were added and shaking was continued for 30 minutes. A further 100  $\mu$ l of 2xYT medium (Amp/Tet) containing the helper phage (1 x 10° cfu/well) was added and shaking was done at RT for 3 hrs. After addition of kanamycin to select for successful helper phage infection, the shaking was continued overnight. The plates were then centrifuged and the supernatants were pipetted directly into ELISA wells coated with 100  $\mu$ I FITC-BSA (100 $\mu$ g/ml) and blocked with milk powder. Washing was performed similarly as during the panning procedure and the bound phages were detected with anti-M13 antibody-POD conjugate (Pharmacia) using soluble POD substrate (Boehringer-Mannheim). Of the 24 clones screened against FITC-BSA, 22 were active in the ELISA (Fig. 13). The initial libraries of similar titer gave rise to no detectable signal.

Specificity for fluorescein was measured in a competitive ELISA. Periplasmic fractions of five FITC specific scFvs were prepared as described above. Western blotting indicated that all clones expressed about the same amount of scFv fragment

(data not shown). ELISA was performed as described above, but additionally, the periplasmic fractions were incubated 30 min at RT either with buffer (no inhibition), with 10 mg/ml BSA (inhibition with BSA) or with 10 mg/ml fluorescein (inhibition with fluorescein) before adding to the well. Binding scFv fragment was detected using the anti-FLAG antibody M1. The ELISA signal could only be inhibited, when soluble fluorescein was added, indicating binding of the scFvs was specific for fluorescein (Fig. 14).

#### 2.6 Sequence analysis

The heavy chain CDR3 region of 20 clones were sequenced in order to estimate the sequence diversity of fluorescein binding antibodies in the library (Fig. 15). In total, 16 of 20 sequences (80%) were different, showing that the constructed library contained a highly diverse repertoire of fluorescein binders. The CDR3s showed no particular sequence homology, but contained on average 4 arginine residues. This bias towards arginine in fluorescein binding antibodies had already been described by Barbas et al., 1992.

#### 2.7 Production

E. coli JM83 was transformed with phagemid DNA of 3 selected clones and cultured in 0.5 L 2xYT medium. Induction was carried out with 1 mM IPTG at OD<sub>600nm</sub> = 0.4 and growth was continued with vigorous shaking at RT overnight. The cells were harvested and pellets were suspended in PBS buffer and sonicated. The supernatants were separated from the cell debris via centrifugation and purified via the BioLogic system (Bio-Rad) by with a POROS MC 20 column (IMAC, PerSeptive Biosystems, Inc.) coupled with an ion-exchange chromatography column. The ion-exchange column was one of the POROS®HS, CM or HQ or PI 20 (PerSeptive Biosystems, Inc.) depended on the theoretical pl of the scFv being purified. The pH of all the buffers was adjusted to one unit lower or higher than the pl of the scFv being purified throughout. The sample was loaded onto the first IMAC column, washed with 7 column volumes of 20 mM sodium phosphate, 1 M NaCl and 10 mM imidazole. This washing was followed by 7 column volumes of 20 mM sodium phosphate and 10 mM imidazole. Then 3 column volumes of an imidazole gradient (10 to 250 mM) were applied and the eluent was connected directly to the ion-exchanger. Nine column volumes of isocratic washing with 250 mM imidazole was followed by 15 column volumes of 250 mM to 100 mM and 7 column volumes of an imidazole / NaCl gradient (100 to 10 mM imidazole, 0 to 1 M NaCl). The flow rate was 5 ml/min. The purity of scFv fragments was checked by SDS-PAGE Coomassie

staining (Fig. 16). The concentration of the fragments was determined from the absorbance at 280 nm using the theoretically determined extinction coefficient (Gill & von Hippel, 1989). The scFv fragments could be purified to homogeneity (see Fig. 16). The yield of purified fragments ranged from 5 to 10 mg/L/OD.

## Example 3: HuCAL H3k2 Library Against a Collection of Antigens

In order to test the library used in Example 2 further, a new selection procedure was carried out using a variety of antigens comprising B-estradiol, testosterone, Lewis-Y epitope (LeY), interleukin-2 (IL-2), lymphotoxin-B (LT-B), E-selectin ligand-1 (ESL-1), and BSA.

#### 3.1 Biopanning

The library and all procedures were identical to those described in Example 2. The ELISA plates were coated with  $\beta$ -estradiol-BSA (100  $\mu$ g/ml), testosterone-BSA (100  $\mu$ g/ml), LeY-BSA (20  $\mu$ g/ml) IL-2 (20  $\mu$ g/ml), ESL-1 (20  $\mu$ g/ml) and BSA (100  $\mu$ g/ml), LT- $\beta$  (denatured protein, 20  $\mu$ g/ml). In the first two rounds, bound phages were eluted with 0.1 M triethylamine (TEA) at RT for 10 minutes. In the case of BSA, elution after three rounds of panning was carried out with addition of BSA in a concentration of 100  $\mu$ g/ml in PBS. In the case of the other antigens, third round elution was done with 0.1 M triethylamine. In all cases except LeY, enrichment of binding phages could be seen (Figure 17). Moreover, a repetition of the biopanning experiment using only the 15-mer library resulted in the enrichment of LeY-binding phages as well (data not shown).

#### 3.2. ELISA measurements

Clones binding to ß-estradiol, testosterone, LeY, LT-ß, ESL-1 and BSA were further analyzed and characterized as described in Example 2 for FITC. ELISA data for anti-ß-estradiol and anti-ESL-1 antibodies are shown in Fig. 18. In one experiment, selectivity and cross-reactivity of binding scFv fragments were tested. For this purpose, an ELISA plate was coated with FITC, testosterone, ß-estradiol, BSA, and ESL-1, with 5 wells for each antigen arranged in 5 rows, and 5 antibodies, one against each of the antigens, were screened against each of the antigens. Fig. 19

shows the specific binding of the antibodies to the antigen it was selected for, and the low cross-reactivity with the other four antigens.

#### 3.3 Sequence analysis

The sequencing data of several clones against ß-estradiol (34 clones), testosterone (12 clones), LT-ß (23 clones), ESL-1 (34 clones), and BSA (10 clones) are given in Figures 20 to 24.

#### **Example 4: Vector Construction**

To be able to take advantage of the modularity of the consensus gene repertoire, a vector system had to be constructed which could be used in phage display screening of HuCAL libraries and subsequent optimization procedures. Therefore, all necessary vector elements such as origins of single-stranded or double-stranded replication, promotor/operator, repressor or terminator elements, resistance genes, potential recombination sites, gene III for display on filamentous phages, signal sequences, or detection tags had to be made compatible with the restriction site pattern of the modular consensus genes. Figure 25 shows a schematic representation of the pCAL vector system and the arrangement of vector modules and restriction sites therein. Figure 25a shows a list of all restriction sites which are already incorporated into the consensus genes or the vector elements as part of the modular system or which are not yet present in the whole system. The latter could be used in a later stage for the introduction of or within new modules.

#### 4.1 Vector modules

A series of vector modules was constructed where the restriction sites flanking the gene sub-elements of the HuCAL genes were removed, the vector modules themselves being flanked by unique restriction sites. These modules were constructed either by gene synthesis or by mutagenesis of templates. Mutagenesis was done by add-on PCR, by site-directed mutagenesis (Kunkel et al., 1991) or multisite oligonucleotide-mediated mutagenesis (Sutherland et al., 1995; Perlak, 1990) using a PCR-based assembly method.

Figure 26 contains a list of the modules constructed. Instead of the terminator module M9 (HindIII-Ipp-PacI), a larger cassette M9II was prepared to introduce Fsel as additional restriction site. M9II can be cloned via HindIII/BsrGI.

All vector modules were characterized by restriction analysis and sequencing. In the case of module M11-II, sequencing of the module revealed a two-base difference in positions 164/65 compared to the sequence database of the template. These two different bases (CA → GC) created an additional BanII site. Since the same two-base difference occurs in the f1 origin of other bacteriophages, it can be assumed that the two-base difference was present in the template and not created by mutagenesis during cloning. This BanII site was removed by site-directed mutagenesis, leading to module M11-III. The BssSI site of module M14 could initially not be removed without impact on the function of the CoIE1 origin, therefore M14-Ext2 was used for cloning of the first pCAL vector series. Figures 29 to 34 are showing the functional maps and sequences of the modules used for assembly of the modular vector pCAL4 (see below). The functional maps and sequences of additional modules can be found in Figure 35a. Figure 35b contains a list of oligonucleotides and primers used for the synthesis of the modules.

### 4.2 Cloning vector pMCS

To be able to assemble the individual vector modules, a cloning vector pMCS containing a specific multi-cloning site (MCS) was constructed. First, an MCS cassette (Fig. 27) was made by gene synthesis. This cassette contains all those restriction sites in the order necessary for the sequential introduction of all vector modules and can be cloned via the 5'-Hindll site and a four base overhang at the 3'-end compatible with an Aatll site. The vector pMCS (Figure 28) was constructed by digesting pUC19 with Aatll and Hindll, isolating the 2174 base pair fragment containing the bla gene and the CoIE1 origin, and ligating the MCS cassette.

# 4.3 Cloning of modular vector pCAL4

This was cloned step by step by restriction digest of pMCS and subsequent ligation of the modules M1 (via Aatll/Xbal), M7III (via EcoRl/HindIII), and M9II (via HindIII/BsrGI), and M11-II (via BsrGI/NheI). Finally, the bla gene was replaced by the cat gene module M17 (via Aatll/BglII), and the wild type CoIE1 origin by module M14-Ext2 (via BglII/NheI). Figure 35 is showing the functional map and the sequence of pCAL4.

# 4.4 Cloning of low-copy number plasmid vectors pCALO

A series of low-copy number plasmid vectors was constructed in a similar way using the p15A module M12 instead of the ColE1 module M14-Ext2. Figure 35a is showing the functional maps and sequences of the vectors pCALO1 to pCALO3.

### Example 5: Construction of a HuCAL scFv Library

#### 5.1. Cloning of all 49 HuCAL scFv fragments

All 49 combinations of the 7 HuCAL-VH and 7 HuCAL-VL consensus genes were assembled as described for the HuCAL VH3-V $\kappa$ 2 scFv in Example 2 and inserted into the vector pBS12, a modified version of the pLisc series of antibody expression vectors (Skerra et *al.*, 1991).

### 5.2 Construction of a CDR cloning cassette

For replacement of CDRs, a universal ß-lactamase cloning cassette was constructed having a multi-cloning site at the 5'-end as well as at the 3'-end. The 5'-multi-cloning site comprises all restriction sites adjacent to the 5'-end of the HuCAL VH and VL CDRs, the 3'-multi-cloning site comprises all restriction sites adjacent to the 3' end of the HuCAL VH and VL CDRs. Both 5'- and 3'-multi-cloning site were prepared as cassettes via add-on PCR using synthetic oligonucleotides as 5'- and 3'-primers using wild type ß-lactamase gene as template. Figure 36 shows the functional map and the sequence of the cassette bla-MCS.

# 5.3. Preparation of VL-CDR3 library cassettes

The VL-CDR3 libraries comprising 7 random positions were generated from the PCR fragments using oligonucleotide templates  $V\kappa 1\&V\kappa 3$ ,  $V\kappa 2$  and  $V\kappa 4$  and primers O\_K3L\_5 and O\_K3L\_3 (Fig. 37) for the  $V\kappa$  genes, and  $V\lambda$  and primers O\_L3L\_5 (5'-GCAGAAGGCGAACGTCC-3') and O\_L3LA\_3 (Fig. 38) for the  $V\lambda$  genes. Construction of the cassettes was performed as described in Example 2.3.

## 5.4 Cloning of HuCAL scFv genes with VL-CDR3 libraries

Each of the 49 single-chains was subcloned into pCAL4 via Xbal/EcoRI and the VL-CDR3 replaced by the β-lactamase cloning cassette via Bbsl/MscI, which was then replaced by the corresponding VL-CDR3 library cassette synthesized as described above. This CDR replacement is described in detail in Example 2.3 where the cat gene was used.

# 5.5 Preparation of VH-CDR3 library cassette

The VH-CDR3 libraries were designed and synthesized as described in Example 2.3.

# 5.6 Cloning of HuCAL scFv genes with VL- and VH-CDR3 libraries

Each of the 49 single-chain VL-CDR3 libraries was digested with BssHII/Styl to replace VH-CDR3. The "dummy" cassette digested with BssHII/Styl was inserted, and was then replaced by a corresponding VH-CDR3 library cassette synthesized as described above.

### **Example 6: Expression tests**

Expression and toxicity studies were performed using the scFv format VH-linker-VL. All 49 combinations of the 7 HuCAL-VH and 7 HuCAL-VL consensus genes assembled as described in Example 5 were inserted into the vector pBS13, a modified version of the pLisc series of antibody expression vectors (Skerra et al., 1991). A map of this vector is shown in Fig. 39.

*E. coli* JM83 was transformed 49 times with each of the vectors and stored as glycerol stock. Between 4 and 6 clones were tested simultaneously, always including the clone H3κ2, which was used as internal control throughout. As additional control, the McPC603 scFv fragment (Knappik & Plückthun, 1995) in pBS13 was expressed under identical conditions. Two days before the expression test was performed, the clones were cultivated on LB plates containing 30  $\mu$ g/ml chloramphenicol and 60 mM glucose. Using this plates an 3 ml culture (LB medium

containing 90 µg chloramphenicol and 60 mM glucose) was inoculated overnight at 37 °C. Next day the overnight culture was used to inoculate 30 ml LB medium containing chloramphenicol (30  $\mu$ g/ml). The starting OD<sub>600nm</sub> was adjusted to 0.2 and a growth temperature of 30 °C was used. The physiology of the cells was monitored by measuring every 30 minutes for 8 to 9 hours the optical density at 600 nm. After the culture reached an  $\mathsf{OD}_{\mathsf{600nm}}$  of 0.5, antibody expression was induced by adding IPTG to a final concentration of 1 mM. A 5 ml aliquot of the culture was removed after 2 h of induction in order to analyze the antibody expression. The cells were lysed and the soluble and insoluble fractions of the crude extract were separated as described in Knappik & Plückthun, 1995. The fractions were assayed by reducing SDS-PAGE with the samples normalized to identical optical densities. After blotting and immunostaining using the α-FLAG antibody M1 as the first antibody (see Ge et al., 1994) and an Fc-specific anti-mouse antiserum conjugated to alkaline phosphatase as the second antibody, the lanes were scanned and the intensities of the bands of the expected size (appr. 30 kDa) were quantified densitometrically and tabulated relative to the control antibody (see Fig. 40).

#### **Example 7: Optimization of Fluorescein Binders**

### 7.1. Construction of L-CDR3 and H-CDR2 library cassettes

A L-CDR3 library cassette was prepared from the oligonucleotide template CDR3L (5'-TGGAAGCTGAAGACGTGGGCGTGTATTATTGCCAGCAG(TR5)(TRI)<sub>4</sub>CCG(TRI)-TTTGGCCAGGGTACGAAAGTT-3') and primer 5'-AACTTTCGTACCCTGGCC-3' for synthesis of the complementary strand, where (TRI) was a trinucleotide mixture representing all amino acids except Cys, (TR5) comprised a trinucleotide mixture representing the 5 codons for Ala, Arg, His, Ser, and Tyr.

A H-CDR2 library cassette was prepared from the oligonucleotide template CDRsH (5'-AGGGTCTCGAGTGGGTGAGC(TRI)ATT(TRI)<sub>2-3</sub>(6)<sub>2</sub>(TRI)ACC(TRI)TATGCGGATA-GCGTGAAAGGCCGTTTTACCATTTCACGTGATAATTCGAAAAACACCCA-3'), and primer 5'-TGGTGTTTTTCGAATTATCA-3' for synthesis of the complementary strand, where (TRI) was a trinucleotide mixture representing all amino acids except Cys, (6) comprised the incorporation of (A/G) (A/C/G) T, resulting in the formation of 6 codons for Ala, Asn, Asp, Gly, Ser, and Thr, and the length distribution being obtained by performing one substoichiometric coupling of the (TRI) mixture during synthesis, omitting the capping step normally used in DNA synthesis.

DNA synthesis was performed on a 40 nmole scale, oligos were dissolved in TE buffer, purified via gel filtration using spin columns (S-200), and the DNA concentration determined by OD measurement at 260 nm (OD 1.0 = 40  $\mu$ g/ml).

10 nmole of the oligonucleotide templates and 12 nmole of the corresponding primers were mixed and annealed at 80°C for 1 min, and slowly cooled down to 37°C within 20 to 30 min. The fill-in reaction was performed for 2 h at 37°C using Klenow polymerase (2.0  $\mu$ l) and 250 nmole of each dNTP. The excess of dNTPs was removed by gel filtration using Nick-Spin columns (Pharmacia), and the double-stranded DNA digested with Bbsl/Mscl (L-CDR3), or Xhol/Sful (H-CDR2) over night at 37°C. The cassettes were purified via Nick-Spin columns (Pharmacia), the concentration determined by OD measurement, and the cassettes aliquoted (15 pmole) for being stored at -80°C.

### 7.2 Library cloning:

DNA was prepared from the collection of FITC binding clones obtained in Example 2 (approx.  $10^4$  to clones). The collection of scFv fragments was isolated via Xbal/EcoRl digest. The vector pCAL4 (100 fmole,  $10~\mu g$ ) described in Example 4.3 was similarly digested with Xbal/EcoRl, gel-purified and ligated with 300 fmole of the scFv fragment collection over night at  $16^{\circ}$ C. The ligation mixture was isopropanol precipitated, air-dried, and the pellets were redissolved in  $100~\mu l$  of dd  $H_2$ O. The ligation mixture was mixed with 1 ml of freshly prepared electrocompetent SCS 101 cells (for optimization of L-CDR3), or XL1 Blue cells (for optimization of H-CDR2) on ice. One round of electroporation was performed and the transformants were eluted in SOC medium, shaken at 37°C for 30 minutes, and an aliquot plated out on LB plates (Amp/Tet/Glucose) at 37°C for 6-9 hrs. The number of transformants was 5 x  $10^4$ .

Vector DNA (100  $\mu$ g) was isolated and digested (sequence and restriction map of scH3 $\kappa$ 2 see Figure 8) with Bbsl/Mscl for optimization of L-CDR3, or Xhol/NspV for optimization of H-CDR2. 10  $\mu$ g of purified vector fragments (5 pmole) were ligated with 15 pmole of the L-CDR3 or H-CDR2 library cassettes over night at 16°C. The ligation mixtures were isopropanol precipitated, air-dried, and the pellets were redissolved in 100  $\mu$ l of dd H<sub>2</sub>O. The ligation mixtures were mixed with 1 ml of freshly prepared electrocompetent XL1 Blue cells on ice. Electroporation was performed and the transformants were eluted in SOC medium and shaken at 37°C for 30 minutes. An aliquot was plated out on LB plates (Amp/Tet/Glucose) at 37°C for 6-9

hrs. The number of transformants (library size) was greater than 10<sup>8</sup> for both libraries. The libraries were stored as glycerol cultures.

### 7.3. Biopanning

This was performed as described for the initial  $H3\kappa2$  H-CDR3 library in Example 2.1. Optimized scFvs binding to FITC could be characterized and analyzed as described in Example 2.2 and 2.3, and further rounds of optimization could be made if necessary.

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Table 1A: Human kappa germline gene segments

Used Name¹	Reference <sup>2</sup>	Family <sup>3</sup>	Germline genes
Vk1-1	9	1	08; 018; DPK1
Vk1-2	1	1	L14; DPK2
Vk1-3	2	1	L15(1); HK101; HK146; HK189
Vk1-4	9	1	L11
Vk1-5	2 .	1	A30
Vk1-6	1 .	1	LFVK5
Vk1-7	1	1	LFVK431
Vk1-8	1.	1	L1; HK137
Vk1-9	1	1	A20; DPK4
Vk1-10	1	1	L18; Va"
Vk1-11	1	1	L4; L18; Va'; V4a
Vk1-12	2	1	L5; L19(1); Vb; Vb4; DPK5; L19(2); Vb"; DPK6
Vk1-13	2	1	L15(2); HK134; HK166; DPK7
Vk1-14	8 .	1 .	L8; Vd; DPK8
Vk1-15	8	1	L9; Ve
Vk1-16	1	1	L12(1); HK102; V1
Vk1-17	2	1	L12(2)
Vk1-18	1.	Ĺ	012a (V3b)
Vk1-19	6	1	O2; O12; DPK9
Vk1-20	2	. 1	L24; Ve"; V13; DPK10
Vk1-21	1	1	04; 014
Vk1-22	2	1	L22
Vk1-23	2	1	L23 -
Vk2-1	1	· 2	A2; DPK12
Vk2-2	6	· 2	01; 011(1); DPK13
Vk2-3	6	2	012(2); V3a
Vk2-4	2	2	L13
Vk2-5	1	2	DPK14
Vk2-6	4	2	A3; A19; DPK15
Vk2-7	4	2	A29; DPK27
Vk2-8	4	2	A13
Vk2-9	1	2	A23

Table 1A: (continued)

Used Name <sup>1</sup>	Reference <sup>2</sup>	Family <sup>3</sup>	Germline genes
Vk2-10	4	2	A7; DPK17
Vk2-11	4	2	A17; DPK18
Vk2-12	4	2	A1; DPK19
Vk3-1	11	3	A11; humkv305; DPK20
Vk3-2	1	3	L20; Vg"
Vk3-3	2	3	L2; L16; humkv328; humkv328h2; humkv328h5; DPK21
Vk3-4	11	· 3	A27; humkv325; VkRF; DPK22
Vk3-5	2	3	L25; DPK23
Vk3-6	2	3	L10(1)
Vk3-7	7	3	L10(2)
Vk3-8	7	3	L6; Vg
Vk4-1	3	4	B3; VkIV; DPK24
Vk5-1	10	5	B2: EV15
Vk6-1	12	6	A14; DPK25
Vk6-2	12	6	A10; A26; DPK26
Vk7-1	5	7	B1

Table 1B: Human lambda germline gene segments

Used Name¹	Reference	Family <sup>3</sup>	Germline genes
DPL1	1	1	
DPL2	1	1 -	HUMLV1L1
DPL3	1	1	HUMLV122
DPL4	1	1	VLAMBDA 1.1
HUMLV117	2	1	
DPL5	1	1	HUMLV117D
DPL6	1 .	1	
DPL7	1	1	IGLV1S2
DPL8	1	1	HUMLV1042
DPL9	1	1	HUMLV101
DPL10	1	2	
VLAMBDA 2.1	3	2	
DPL11	1	Ź	
DPL12	. 1	2	
DPL13	1	2	
DPL14	1	2	
DPL16	1	3	Humlv418; IGLV3S1
DPL23	. 1 · ·	3	VI III.1
Humlv318	4	3	
DPL18	1	7	4A; HUMIGLVA
DPL19	1	7	
DPL21	1	8	VL8.1
HUMLV801	5	8	
DPL22	1	9 .	
DPL24	1	unassigned	VLAMBDA N.2
gVLX-4.4	6	10	•

Table 1C: Human heavy chain germline gene segments

Used Name <sup>1</sup>	Reference <sup>2</sup>	Family <sup>3</sup>	Germline genes
VH1-12-1	19	1	DP10; DA-2; DA-6
VH1-12-8	22	1	RR.VH1:2
VH1-12-2	6	1	hv1263
VH1-12-9	7	1	YAC-7; RR.VH1.1; 1-69
VH1-12-3	19	1	DP3
VH1-12-4	· 19	1	DP21; 4d275a; VH7a
VH1-12-5	18	1	1-4.1b; V1-4.1b
VH1-12-6	21	1	1D37; VH7b; 7-81; YAC-10
VH1-12-7	19	1	DP14; VH1GRR; V1-18
VH1-13-1	10	1	71-5; DP2
VH1-13-2	10	1	E3-10
VH1-13-3	19	1	DP1
VH1-13-4	12	1	V35
VH1-13-5	8	1	V1-2b
VH1-13-6	18	1	I-2; DP75
°VH1-13-7	<b>21</b> .	1	V1-2
VH1-13-8	19	1	DP8
VH1-13-9	3	1	1-1
VH1-13-10	19	1	DP12
VH1-13-11	15	1	V13C
VH1-13-12	18	1	I-3b; DP25; V1-3b
VH1-13-13	3	1	1-92
VH1-13-14	· 18	1	I-3; V1-3
VH1-13-15	19	1	DP15; V1-8
VH1-13-16	3	1	21-2; 3-1; DP7; V1-46
VH1-13-17	16	1	HG3
VH1-13-18	19	. 1	DP4; 7-2; V1-45
VH1-13-19	27	1	COS 5
VH1-1X-1	19	1	DP5: 1-24P
VH2-21-1	18	2	II-5b
VH2-31-1	2	2	VH2S12-1
VH2-31-2	2	2	VH2S12-7
VH2-31-3	2	2	VH2S12-9; DP27
VH2-31-4	2	2	VH2S12-10
VH2-31-5	14	2	V2-26; DP26; 2-26
VH2-31-6	15	2	VF2-26

Table 1C: (continued)

Used Name <sup>1</sup>	Reference <sup>2</sup>	Family <sup>3</sup>	Germline genes
VH2-31-7	19	2	DP28; DA-7
VH2-31-14	7	2	YAC-3; 2-70
VH2-31-8	2	2	VH2S12-5
VH2-31-9	2	2	VH2S12-12
VH2-31-10	18	2	II-5; V2-5
VH2-31-11	2	2	VH2S12-2; VH2S12-8
VH2-31-12	2	2	VH2S12-4; VH2S12-6
VH2-31-13	2 .	2	VH2S12-14
VH3-11-1	13	3	v65-2; DP44
VH3-11-2	19	3	DP45
VH3-11-3	3	3	13-2; DP48
VH3-11-4	19	3	DP52
VH3-11-5	14	3	v3-13
VH3-11-6	19	3	DP42
VH3-11-7	3	3	8-1B; YAC-5; 3-66
VH3-11-8	14	3	V3-53
VH3-13-1	3	3	22-2B; DP35; V3-11
VH3-13-5	19	3	DP59; VH19; V3-35
VH3-13-6	25	3	f1-p1; DP61
VH3-13-7	19	3	DP46; GL-SJ2; COS 8; hv3005; hv3005f3; 3d21b; 56p1
VH3-13-8	24	3	VH26
VH3-13-9	5	3	vh26c
VH3-13-10	19 .	3	DP47; VH26; 3-23
VH3-13-11	3	3	1-91
VH3-13-12	19	3	DP58
VH3-13-13	3	3	1-9III; DP49; 3-30; 3d28.1
VH3-13-14	24	. 3	3019B9; DP50; 3-33; 3d277
VH3-13-15	27	3	COS 3
VH3-13-16	19	3	DP51
VH3-13-17	16	3	H11
VH3-13-18	19	3	DP53; COS 6; 3-74; DA-8
VH3-13-19	19	3	DP54; VH3-11; V3-7
VH3-13-20	14	3	V3-64, YAC-6
VH3-13-21	14	3	V3-48
VH3-13-22	14	3	V3-43; DP33
VH3-13-23	14	3	V3-33

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Table 1C: (continued)

Used Name'	Reference <sup>2</sup>	Family	Germline genes
VH3-13-24	14	3	V3-21; DP77
VH3-13-25	14	3	V3-20; DP32
VH3-13-26	14	3	V3-9; DP31
VH3-14-1	3	3	12-2; DP29; 3-72; DA-3
VH3-14-4	7	. 3	YAC-9; 3-73; MTGL
VH3-14-2	. 4	3	VHD26
VH3-14-3	19	3	DP30
VH3-1X-1	1	3	LSG8.1; LSG9.1; LSG10.1; HUM12IGVH; HUM13IGVH
VH3-1X-2	1	3	LSG11.1; HUM4IGVH
VH3-1X-3	3	3	9-1; DP38; LSG7.1; RCG1.1; LSG1.1; LSG3.1; LSG5.1; HUM15IGVH; HUM2IGVH; HUM9IGVH
VH3-1X-4	1	3	LSG4.1
VH3-1X-5	1	3	LSG2.1
VH3-1X-6	1	3	LSG6.1; HUM10IGVH
VH3-1X-7	18	3	3-15; V3-15
VH3-1X-8	1	<b>3</b> .	LSG12.1; HUM5IGVH
VH3-1X-9	14	3	V3-49
VH4-11-1	22	4	Tou-VH4.21
VH4-11-2	17	4	VH4.21; DP63; VH5; 4d76; V4-34
VH4-11-3	23	4	4.44
VH4-11-4	<b>23</b> .	4	4.44.3
VH4-11-5	23	4	4.36
VH4-11-6	23	4	4.37
VH4-11-7	18	4	IV-4; 4.35; V4-4
VH4-11-8	17	4	VH4.11; 3d197d; DP71; 58p2
VH4-11-9	20	4	H7
VH4-11-10	20	4	H8
VH4-11-11	20	· 4	Н9
VH4-11-12	17	4	VH4.16
VH4-11-13	. 23	4	4.38
VH4-11-14	17	4	VH4.15
VH4-11-15	11	4	58
VH4-11-16	10	4	71-4; V4-59
VH4-21-1	11	4	11
VH4-21-2	17	4	VH4.17; VH4.23; 4d255; 4.40; DP69
VH4-21-3	17	4	VH4.19; 79; V4-4b

Table 1C: (continued)

Used Name'	Reference <sup>2</sup>	Family <sup>3</sup>	Germline genes
VH4-21-4	19	. 4	DP70; 4d68; 4.41
VH4-21-5	19	4	DP67; VH4-4B
VH4-21-6	17	4	VH4.22; VHSP; VH-JA
VH4-21-7	17	4	VH4.13; 1-9II; 12G-1; 3d28d; 4.42; DP68; 4-28
VH4-21-8	26	4	hv4005; 3d24d
VH4-21-9	17	4	VH4.14
VH4-31-1	23	4	4.34; 3d230d; DP78
VH4-31-2	23	4	4.34.2
VH4-31-3	19	4	DP64; 3d216d
VH4-31-4	19	4	DP65; 4-31; 3d277d
VH4-31-5	23	4	4.33; 3d75d
VH4-31-6	20	4	H10
VH4-31-7	20	4	H11
VH4-31-8	23	· 4	4.31
VH4-31-9	23	4	4.32
VH4-31-10	20	. 4	3d277d
VH4-31-11	20	4	3d216d
VH4-31-12	20	4 .	3d279d
VH4-31-13	17	4	VH4.18; 4d154; DP79
VH4-31-14	8	4	V4-39
VH4-31-15	11 .	4	2-1; DP79
VH4-31-16	23	4	4.30
VH4-31-17	17	4	VH4.12
VH4-31-18	10	4	71-2; DP66
VH4-31-19	23	4	4.39
VH4-31-20	8	4	V4-61
VH5-12-1	9	. <b>5</b>	VH251; DP73; VHVCW; 51-R1; VHVLB; VHVCH; VHVIT; VHVAU; VHVBLK; VhAU; V5-51
VH5-12-2	17	5	VHVJB
VH5-12-3	3	5	1-v; DP80; 5-78
VH5-12-4	9 .	5	VH32; VHVRG; VHVMW; 5-2R1
VH6-35-1	4	6	VHVI; VH6; VHVIIS; VHVITE; VHVIJB; VHVICH; VHVICW; VHVIBLK; VHVIMW; DP74; 6-1G1; V6-1

Table 2A: rearranged human kappa sequences

Name <sup>1</sup>	aa²	Computed family <sup>3</sup>	Germline gene <sup>4</sup>	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference'
III-3R	108	1	08	1	1,1%	70
No.86	109	1	08	3	3,2%	80
AU	108	1 .	80	6	6,3%	103
ROY	108	1	80	6	6,3%	43
IC4	108	1	08	6	6,3%	70
HIV-B26	106	1	80	3	3,2%	8
GRI	108	1	08	8	8,4%	30
AG	106	1	08	<b>8</b> .	8,6%	116
REI	108	1	08	9	9,5%	86
CLL PATIENT 16	88	1	08	2	2,3%	122
CLL PATIENT 14	87	1	08	2	2,3%	122
CLL PATIENT 15	88	1	08	. 2	2,3%	122
GM4672	108	1	08	11	11,6%	24
HUM. YFC51.1	108	1	08	12	12,6%	110
LAY	108	1	08	12	12,6%	48
HIV-b13	106	1	08	9	9,7%	8
MAL-NaCl	108	1	08	13	13,7%	102
STRAb SA-1A	108	1	02	0	0,0%	120
HuVHCAMP	108	1	08	13	13,7%	100
CRO	108	1	02	10	10,5%	30
Am107	108	1	02	12	12,6%	108
WALKER	107	1	02	4	4,2%	57
III-2R	109	1	A20	0	0,0%	70
FOG1-A4	107	1	A20	4	4,2%	41
HK137	95	1	L1	. 0	0,0%	10
CEA4-8A	107	1	02	7	7,4%	41
Va'	95	1	L4	0	0,0%	90
TR1.21	108	1	02	4	4,2%	92
HAU	108	1	02	6	6,3%	123
HK102	95	1	L12(1)	0	0,0%	9
H20C3K	108	1	L12(2)	3	3,2%	125
CHEB	108	ĺ	02	7	7,4%	5
HK134	95	1	L15(2)	0	0,0%	10
TEL9	108	1	02	9	9,5%	73

Table 2A: (continued)

Name¹	aa²	Computed family <sup>3</sup>	Germline gene <sup>4</sup>	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference <sup>7</sup>
TR1.32	103	1	02	3	3,2%	92
RF-KES1	97	1	A20	4	4,2%	121
WES	108	1	L5	10	10,5%	61
DILp1	95	1	04	1	1,1%	70
SA-4B	107	1	L12(2)	8	8,4%	120
HK101	95	1	L15(1)	0	0,0%	9
TR1.23	108	1	02	5	5,3%	92
HF2-1/17	108	1	A30	0	0,0%	4
2E7	108	1	A30	1	1,1%	62
33.C9	107	1	L12(2)	7	7,4%	126
3D6	105	. 1	L12(2)	2	2,1%	34
l-2a	108	1	L8	8	8,4%	. ~ 70
RF-KL1	97 ·	1	L8	4	4,2%	121
TNF-E7	108	1	A30	9	9,5%	41
TR1.22	108	1	02	7	7,4%	92
HIV-B35	106	1	02	2	2,2%	8
HIV-b22	106	1	02	. 2	2,2%	8
HIV-b27	106	1	02	2	2,2%	8
HIV-B8	107	1	02	10	10,8%	8
HIV-b8	107	1	02	10	10,8%	8
RF-SJ5	95	1 .	A30	5	5,3%	113
GAL(I)	108	1	A30	6	6,3%	64
R3.5H5G	108	1	02	6	6,3%	70
HIV-b14	106	1	A20	2	2,2%	8
TNF-E1	105	1	·L5	8	8,4%	41
WEA	108	1	A30	8	8,4%	37
EU	108	1	L12(2)	5	5,3%	40
FOG1-G8	108	1	L8	11	11,6%	41
1X7RG1	108	1	L1	8	8,4%	70
BLI	108	1	L8	3	3,2%	72
KUE	108	1	L12(2)	11	11,6%	32
LUNm01	108	1	L12(2)	10	10,5%	6
HIV-b1	106	1	A20	4	4,3%	8
HIV-s4	103	1	02	2	2,2%	8

Table 2A: (continued)

Name <sup>1</sup>	aa²	Computed family <sup>3</sup>	Germline gene <sup>4</sup>	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference <sup>7</sup>
CAR	107	1	L12(2)	11	11,7%	79
BR	- 107	1	L12(2)	11	11,6%	50
CLL PATIENT 10	88	1	02	0	0,0%	122
CLL PATIENT 12	88	. 1	02	0	0,0%	122
KING	108	1 .	L12(2)	12	12,6%	30
V13	95	1	L24	0	0,0%	46
CLL PATIENT 11	87	1	02	0	0,0%	122
CLL PATIENT 13	87	1	02	0	0,0%	122
CLL PATIENT 9	. 88	1	012	1	1,1%	122
HIV-B2	106	1	A20	9	9,7%	8
HIV-b2	106	1	A20	9	9,7%	8
CLL PATIENT 5	88	1	A20	1	1,1%	122
CLL PATIENT 1	88	· 1	L8	2	2,3%	122
CLL PATIENT 2	88	1	L8	0	0,0%	122
CLL PATIENT 7	88	1	L5	0	0,0%	122
CLL PATIENT 8	. 88	1	L5	0	0,0%	122
HIV-b5	105	1 .	L5	11	12,0%	8
CLL PATIENT 3	87	. 1	L8	1	1,1%	122
CLL PATIENT 4	88	1	L9	0	0,0%	122
CLL PATIENT 18	85	1	L9	6	7,1%	122
CLL PATIENT 17	86	1	L12(2)	7	8,1%	122
HIV-b20	107	3	A27	11	11,7%	8
2C12	108	1 '	L12(2)	20	. 21,1%	68
1B11	108	1	L12(2)	20	21,1%	68
1H1	108	1	L12(2)	21	22,1%	68
2A12	108	1	L12(2)	21	22,1%	68
CUR	109	3	A27	0	0,0%	66
GLO	109	3	A27	0	0,0%	16
RF-TS1	96	3	A27	0	0,0%	121
GAR'	109	3	A27	0	0.0%	67
FLO	109	3	A27	0	0,0%	66
PIE	109	3	A27	0	0.0%	91
HAH 14.1	109	3	A27	1 .	1,0%	51
HAH 14.2	109	3	A27	1	1,0%	51

Table 2A: (continued)

Name <sup>1</sup>	aa²	Computed family <sup>3</sup>	Germline gene⁴	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference <sup>7</sup>
HAH 16.1	109	3	A27	. 1	1,0%	51
NOV .	109	3	A27	17.	1,0%	52
33.F12	108	3	A27	1	1,0%	126
8E10	110	3	A27	1	1,0%	25
тнз	109	3	A27	1	1,0%	25
HIC (R)	108	3	A27	0	0,0%	51
SON	110	3	A27	1	1,0%	<b>67</b> .
PAY	109	3	A27	1	1,0%	66
GOT	109	3	A27	1	1,0%	67
mAbA6H4C5	109	3	A27	1	1,0%	12
BOR'	109	3	A27	2	2,1%	84
RF-SJ3	96	3	A27	2 -	2,1%	121
SIE .	109	3	A27	2	2,1%	15
ESC	109	3	A27	2	2,1%	98
HEW'	110	3	A27	2	2,1%	98
YES8c	109	3	A27	3	3,1%	33
TI	109	3	A27	3	3,1%	114
mAb113	109	3	A27	3	3,1%	. 71
HEW	107	3	A27	0	0,0%	94
BRO	106	3	A27	0	0,0%	94
ROB	106	3	A27	. 0	0,0%	94
NG9	96	3	A27	4	4,2%	11
NEU	109	3	A27	4	4,2%	66
WOL	109	3	A27	4	4,2%	2
35G6	109	3	A27	4	4,2%	59
RF-SJ4	109	3	A11	0	0.0%	88
KAS 🕠	109	3	A27	4	4,2%	84
BRA	106	3	A27	1	1,1%	94
HAH	106	3	A27	1	1,1%	94
HIC	105	3	A27	0	0,0%	94
FS-2	109	. 3	A27	6	6,3%	87
JH'	107	3	A27	6	6,3%	38
EV1-15	109	3	A27	<b>6</b> .	6,3%	83
SCA	108	3	A27	6	6,3%	65
			56			

Table 2A: (continued)

Name <sup>1</sup>	aa²	Computed family <sup>3</sup>	Germline gene⁴	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference <sup>7</sup>
mAb112	109	3	A27	6	6,3%	71
SIC .	103	3	A27	3	3,3%	94
SA-4A	109	3	A27	6	6,3%	120
SER	108	3	A27	6	6,3%	98
GOL'	109	3	A27	7	7,3%	82
B5G10K	105	3	A27	9	9,7%	125
HG2B10K	110	3	A27	<del>_9</del>	9,4%	125
Taykv322	105	3	A27	5	5,4%	52
CLL PATIENT 24	89	3	A27	1	1,1%	122
HIV-b24	107	3	A27	7	7,4%	8
HIV-b6	107	3	A27	7	7,4%	8
Taykv310	99	3	A27	1	1,1%	52
KA3D1	108	3	L6	0	0,0%	85
19.E7	107	3	L6	. 0	0,0%	126
rsv6L	109	3	A27	12	12,5%	7
Taykv320	98	3	A27	1	1,2%	52
Vh	96	3	L10(2)	0	0,0%	89
LS8	108	3	L6	1	1,1%	109
LS1	108	3	L6	1	1,1%	109
LS2S3-3	107	3	L6	2.	2,1%	99
LS2	· 108	3	L6	1,	1,1%	109
LS7	108	3	L6	1	1,1%	109
LS2S3-4d	107	3	L6	2	2,1%	99
LS2S3-4a	107	3	L6	. 2	2,1%	99
LS4	108	3	L6	1	1,1%	109
LS6	108	3	L6	1	1,1%	109
LS2S3-10a	107	3	L6	2	2,1%	99
LS2S3-8c	107	3	L6	2	2,1%	99
LS5	108	3	L6	1	1,1%	109
LS2S3-5	107	3	L6	3	3,2%	99
LUNm03	109	3	A27	13	13,5%	6
IARC/BL41	108	3	A27	13	13,7%	55
slkv22	99	3	A27	3	3,5%	13
POP	108	3	L6	4	4,2%	111

Table 2A: (continued)

Name <sup>1</sup>	aa²	Computed family <sup>3</sup>	Germline gene <sup>4</sup>	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference <sup>7</sup>
LS2S3-10b	107	3	L6	3	3,2%	99
LS2S3-8f	107	3	L6	3	3,2%	99
LS2S3-12	107	3	L6	3	3,2%	99
HIV-B30	107	3	A27	11	11,7%	8
HIV-B20	107	3	A27	11	11,7%	8
HIV-b3	108	3	A27	11	11,7%	8
HIV-s6	104	3	A27	9	9,9%	8
YSE	107	3	L2/L16	1	1,1%	72
POM	109	3	L2/L16	9	9,4%	53
Humkv328	95	3	L2/L16	1	1,1%	19
CLL	109	3	L2/L16	3	3,2%	47
LES	96	3	L2/L16	3	3,2%	38
HIV-s5	104	3	A27	11	12,1%	8
HIV-s7	104	3	<b>A27</b>	11	12,1%	8
slkv1	99	3	A27	7	8,1%	13
Humka31es	95	3	L2/L16	4	4,2%	18
slkv12	101	3	A27	8	9,2%	13
RF-TS2	95	3	L2/L16	<b>3</b> ·	3,2%	121
II-1	109	3	L2/L16	4	4,2%	70
HIV-s3	105	3	A27	13	14,3%	8
RF-TMC1	96	3 .	<b>L6</b>	10	10,5%	121
GER	109	. 3	L2/L16	7 .	7,4%	75 ·
GF4/1.1	109	3	L2/L16	8	8,4%	36
mAb114	109	3	L2/L16	6	6,3%	71
HIV-loop13	109	3	L2/L16	7	7,4%	8
bkv16	86	3	L6	1	1,2%	13
CLL PATIENT 29	86	3	L6	1	1,2%	122
slkv9	98	3	L6	3	3,5%	13
bkv17	99	3	L6	1	1,2%	13
slkv14	99	3	L6	1	1,2%	13
slkv16	101	3	L6	2	2,3%	13
bkv33	101	3	L6	4	4,7%	13
slkv15	99	- 3	L6	2	2,3%	13
bkv6	100	3	L6	3	3,5%	13

Table 2A: (continued)

Name <sup>1</sup>	aa²	Computed family <sup>3</sup>	Germline gene⁴	Diff. to germline <sup>s</sup>	% diff. to germline <sup>6</sup>	Reference <sup>2</sup>
R6B8K	108	3	L2/L16	12	12,6%	125
AL 700	107	3	L2/L16	9	9,5%	117
slkv11	100	3	L2/L16	3	3,5%	13
slkv4	97	3 ·	L6	4	4.8%	13
CLL PATIENT 26	87	3	L2/L16	1	1,1%	122
AL Se124	103	3	L2/L16	9	9,5%	117
slkv13	100	3	L2/L16	6	7,0%	.13
bkv7	100	3	L2/L16	5	5,8%	13
bkv22	100	3	L2/L16	6	7,0%	13
CLL PATIENT 27	84	3	L2/L16	0	0,0%	122
bkv35	100	3	L6	8	9,3%	13
CLL PATIENT 25	87	3	L2/L16	4	4,6%	122
slkv3	86	3	L2/L16	7	8,1%	13
slkv7	99	. 1	02	7	8,1%	13
HuFd79	111	3	L2/L16	24	24,2%	21
RAD	99	3	A27	9	10,3%	78
CLL PATIENT 28	83	3	L2/L16	4	4,8%	122
REE	104	3	L2/L16	25	27,2%	95
FR4	99	3	A27	8	9,2%	77
MD3.3	92	· 3	L6	1	1,3%	54
MD3.1	92	3	L6	0	0,0%	54
GA3.6	92	3	L6	· <b>2</b>	2,6%	54
M3.5N	92	3	L6	3	3,8%	54
MEI,	82	3	A27	0	0,0%	65
MD3.4	92	3	L2/L16	1	1,3%	54
MD3.2	91	3	L6	3	3,8%	54
VER	97	3	A27	19	22,4%	20
CLL PATIENT 30	78	3	L6	3	3,8%	122
M3.1N	92	3	L2/L16	1	1,3%	54
MD3.6	91	3	L2/L16	0	0,0%	54
MD3.8	91	3	L2/L16	0	0.0%	. 54
GA3.4	92	3	L6	7	9,0%	54
M3.6N	92	. 3	A27	0	0,0%	54
	92	3	A27			

Table 2A:

(continued)

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Name <sup>1</sup>	aa²	Computed family <sup>3</sup>	Germline gene⁴	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference <sup>1</sup>
MD3.13	91	3	A27	0	0,0%	54
MD3.7	93	3	A27	0	0,0%	54
MD3.9	93	3	A27	0 .	0,0%	54
GA3.1	93	3	A27	6	7,6%	54
bkv32	101	3	A27	5	5,7%	13
GA3.5	93	3	A27	5	6,3%	54
GA3.7	92	3	A27	_7	8,9%	54
MD3.12	92	3	A27	2	2,5%	54
M3.2N	90	3	L6	. 6	7,8%	54
MD3.5	92	3	A27	1	1,3%	54
M3.4N	91	. 3	L2/L16	8	10,3%	54
M3.8N	91	3	L2/L16	7	9,0%	54
M3.7N	92	3	A27	3	3,8%	54
GA3.2	92	. 3	A27	9	11,4%	54
GA3.8	93	3	A27	4	5,1%	54
GA3.3	92	3	A27	8	10,1%	54
M3.3N	92	3	A27	5	6,3%	54
B6	83	. 3	A27	8	11,3%	78
E29.1 KAPPA	<b>78</b>	3	L2/L16	0	0,0%	22
SCW	108	1	08	12	12,6%	31
REI-based CAMPATH-9	107	1	08	14	14,7%	39
RZ	107	1	08	14	14,7%	50
BI	108	1	08	14	14,7%	14
AND	107	1	02	13	13,7%	69
2A4	109	1	02	12	12,6%	23
KA	108	1	08	19	20,0%	107
MEV	109	1	02	14	14,7%	29
DEE	106	1	02	13	14,0%	76
OU(IOC)	108	1	02	18	18,9%	60
HuRSV19VK	111	1	08	21	21,0%	115
SP2	108	1	02	17	17,9%	93
BJ26	99	1 .	08	21	24,1%	1 .
NI ·	112	1	08	24	24,2%	106
BMA 0310EUCIV2	106	1	L12(1)	21	22,3%	105

Table 2A: (continued)

Name <sup>1</sup>	aa²	Computed family <sup>3</sup>	Germline gene⁴	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference <sup>7</sup>
CLL PATIENT 6	71	1	A20	0	0,0%	122
BJ19	85	1	08	16	21,9%	· 1
GM 607	113	2	A3	0	0,0%	58
R5A3K	- 114	2	A3	1	1,0%	125
R1C8K	114	2	<b>A3</b>	1	1,0%	125
VK2.R149	113	2	A3	. 2	2,0%	118
TR1.6	109	2	<b>A3</b>	4	4,0%	92
TR1.37	104	2	<b>A3</b>	5	5,0%	92
FS-1	113	2	<b>A</b> 3	6	6,0%	87
TR1.8	110	2	<b>A</b> 3	6	6,0%	92
NIM	113	2	<b>A</b> 3	. 8	8,0%	28
Inc	112	2	<b>A</b> 3	11	11,0%	35
TEW	107	2	A3	6	6,4%	96
CUM	114	2	01	7	6.9%	44
HRF1	71	2	<b>A</b> 3	4	5,6%	124
CLL PATIENT 19	87	2	<b>A3</b>	0	0,0%	122
CLL PATIENT 20	87	2	A3	0	0,0%	122
MIL	112	2	<b>A</b> 3	16	16,2%	26
FR	113	2	A3	20	20,0%	101
MAL-Urine	83	1	02	6	8,6%	102
Taykv306	73	3	A27	1	1,6%	52
Taykv312	75	3	A27	1	1,6%	<b>52</b> .
HIV-b29	93	3	A27	14	17,5%	. 8
1-185-37	110	3	A27	. 0	0,0%	119
1-187-29	110	3	A27	0	0,0%	119
Π117	110	.3	A27	9	9,4%	. 63
HIV-loop8	108	3	A27	16	16,8%	8
rsv23L	108	3	A27	16	16,8%	7
HIV-b7	107	3	A27	14	14,9%	8
HIV-b11	107	3	A27	15	16,0%	8
HIV-LC1	107	3	A27	19	20,2%	8
HIV-LC7	107	3	A27	20	21,3%	8
HIV-LC22	107	3	A27	21	22,3%	8
HIV-LC13	107	3	A27	21	22,3%	8

Table 2A: (continued)

Name¹	aa²	Computed family <sup>3</sup>	Germline gene <sup>4</sup>	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference'
HIV-LC3	107	3	A27	21	22,3%	8
HIV-LC5	107	3	A27	21	22,3%	8
HIV-LC28	107	3	A27	21	22,3%	. 8
HIV-b4	107	3	A27	22	23,4%	8
CLL PATIENT 31	87	3	A27	15	17,2%	122
HIV-loop2	108	3	L2/L16	17	17,9%	8
HIV-loop35	108	3 .	L2/L16	17	17,9%	8
HIV-LC11	107	3	A27	23	24,5%	8
HIV-LC24	107	3	A27	23	24,5%	8
HIV-b12	107	3	A27	24	25,5%	8
HIV-LC25	107	3	A27	24	25,5%	8
HIV-b21	107	3	A27	24	25,5%	8
HIV-LC26	107	3	A27	26	27,7%	8.
G3D10K	108	1	L12(2)	12	12,6%	125
Π125	108	1	L5	8	8,4%	63
HIV-s2	103	3	A27	28	31,1%	8
265-695	108	1	L5	7	7,4%	3
2-115-19	108	. 1	A30	2	2,1%	119
rsv13L	107	1	02	20	21,1%	7
HIV-b18	106	1	02	14	15,1%	8
RF-KL5	98	3	L6	36	36,7%	97
ZM1-1	113	2	A17	7	7,0%	3
HIV-s8	103	1	08	16	17,8%	8
K- EV15	95	5	B2	0	0,0%	112
RF-TS3	100	2	A23	0	0,0%	121
HF-21/28	111	2	A17	1	1,0%	17
RPMI6410	113	2	A17	1	1,0%	42
JC11	113	2	A17	1	1,0%	49
0-81	114	2	A17 .	5	5,0%	45
FK-001	113	4	В3	0	0,0%	81
CD5+.28	101	4	В3	1	1,0%	27
LEN	114	4	В3	1	1,0%	104
UC	114	4	В3	1	1,0%	- 111
CD5+.5	101	4	В3	1	1,0%	27

Table 2A: (continued)

Name <sup>1</sup>	aa²	Computed family <sup>3</sup>	Germline gene⁴	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference'
CD5+.26	101	4	В3	1	1,0%	27
CD5+.12	101	4	В3	2	2,0%	27
CD5+.23	101	4	В3	2	2,0%	27
CD5+.7	101	4	В3	2	2.0%	27
VJI	113	4	В3	3	3,0%	56
LOC ·	113	4	В3	3	3,0%	72
MAL	113	4	В3	3	3,0%	72
CD5+.6	101	4	<b>B</b> 3	3	3,0%	27
H2F	113	4	В3	<b>3</b> .	3,0%	70
PB17IV	114	4	В3	4	4,0%	74
CD5+.27	101	4	В3	4	4,0%	27
CD5+.9	101	4	В3	4	4,0%	27 ·
CD528	101	4	B3	5	5,0%	27
CD526	101	4	В3	6	5,9%	27
CD5+.24	101	4	В3	6	5,9%	27
CD5+.10	101	4	В3	6	5,9%	27
CD519	101	4	B3	6	5,9%	27
CD518	101	4	В3	7	6,9%	27
CD516	101	4	В3	8	7,9%	27
CD524	101	4	В3	8	7,9%	27
CD517	101	4	B3	10	9,9%	27
MD4.i	92	4	· B3	0	0,0%	54
MD4.4	92	4	В3	0	0,0%	<b>54</b> ,
MD4.5	92	4	В3	0	0,0%	54 ુ
MD4.6	92	4	В3	0	0,0%	54
MD4.7	92	4	В3	0	0,0%	54
MD4.2	92	4	В3	1	1,3%	54
MD4.3	92	4	В3	5	6,3%	54
CLL PATIENT 22	87	2	A17	2	2,3%	122
CLL PATIENT 23	84	2	A17	2	2,4%	122

Table 2B: rearranged human lambda sequences

Name¹	aa²	Computed family <sup>3</sup>	Germline gene⁴	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference
WAH	110	1	DPL3	7	7%	68
1B9/F2	112	1	DPL3	7	7%	9
DIA	112	1	DPL2	7	7%	36
mAb67	- 89	1	DPL3	0	0%	29
HiH2	110	1	DPL3	12	11%	3
NIG-77	. 112	1	DPL2	. 9	9%	72
OKA	112	1	DPL2	7	7%	84
KOL	112	1	DPL2	12	11%	40
T2:C5	111	1	DPL5	0	0%	6
T2:C14	110	. 1	DPL5	0 .	0%	6
PR-TS1	110	1	DPL5	0	0%	55
4G12	111	1	DPL5	1	1%	35
KIM46L	112	1	HUMLV117	0	0%	8
Fog-B	111	1	DPL5	3	3%	31
9F2L	111	1	DPL5	3	3%	79
mAb111	110	1	DPL5	3	3%	48
PHOX15	111	1	DPL5	4	4%	49
BL2	111	1	DPL5	4	4%	74
NIG-64	111	, 1	DPL5	4	4%	72
RF-SJ2	100	1	DPL5	6	6%	78
AL EZI	112	1	· DPL5	7	<b>7%</b>	41
ZIM	112	. 1	HUMLV117	7	7%	18
RF-SJ1	100	1.	DPL5	9	9%	78
IGLV1.1	98	1	DPL4	0	0%	. 1
NEW	112	1	HUMLV117	11	10%	42
CB-201	87	. 1	DPL2	1	1%	62
MEM	109	1	DPL2	6	6%	50
H210	111	. 2	DPL10	4	4%	45
NOV	110	2	DPL10	8	8%	25
NEI	111	2	DPL10	8	8%	24
AL MC	110	2	DPL11	6	6%	28
MES	112	2	DPL11	8	8%	84
FOG1-A3	. 111	2	DPL11	9 .	. 9%	27
AL NOV	112	2	DPL11	7	7%	28

Table 2B: (continued)

Name¹	aa²	Computed family <sup>3</sup>	Germline gene⁴	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference <sup>7</sup>
HMST-1	110	2	DPL11	4	4%	82
HBW4-1	108	2	DPL12	9	9%	52
wh.	110	2	DPL11	11	11%	34
11-50	110	2	DPL11	7	70/0	82
НВр2	110	2	DPL12	8	8%	. 3
NIG-84	113	2	DPL11	12	11%	73
VIL	112	2	DPL11	9	9%	58
TRO	111	2	DPL12	10	10%	61
ES492	108	2	DPL11	15	15%	76
mAb216	89	2	DPL12	1	1%	7
BSA3	109	3	DPL16	0	0%	49
THY-29	110	3	DPL16	0 -	0%	27
PR-TS2	108	3	DPL16	0	0%	55
E29.1 LAMBDA	107	3	DPL16	1	10/0	13
mAb63	109	3	DPL16	2	2%	29
TEL14	110	3	DPL16	6	6%	49
6H-3C4	108	3	DPL16	7	7%	39
SH	109	3	DPL16	7	7%	70
AL GIL	109	3	DPL16	8	8%	23
H6-3C4	108	3	DPL16	8	8%	83
V-lambda-2.DS	111	2	DPL11	3	3%	15
8.12 ID	110	2	DPL11	3	3%	81
DSC	111	2	DPL11	3	3%	56
PV11	110	. 2	DPL11	1	1%	56
33.H11	110	2	DPL11	4	4%	81
AS17	111	2	DPL11	7	7%	56
SD6	110	2	DPL11	7	7%	56
KS3	110	2	DPL11	9	9%	56
PV6	110	2	DPL12	5	5%	. 56
NGD9	110	2	DPL11	7	7%	56
MUC1-1	111	2	DPL11	11	.10%	27
A30c	111	2	DPL10	6	6%	56
KS6	110	2	DPL12	6	6%	56
TEL13	111	2	DPL11 65	11	10%	49

Table 2B: (continued)

Name¹	aa²	Computed family <sup>3</sup>	Germline gene <sup>4</sup>	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference
AS7	110	2	DPL12	6	6%	56
MCG	112	2	DPL12	12	11%	20
U266L	110	2	DPL12	13	12% ·	77
PR-SJ2	110	2	DPL12	14	13%	55
вон	112	2	DPL12	11	10%	37
TOG	111	2	DPL11	19	18%	53
TEL16	111	2	DPL11	19	18%	49
No.13	110	2	DPL10	14	13%	.52
ВО	112	2	DPL12	18	17%	80
WIN	112	2	DPL12	17	16%	11
BUR	104	2	DPL12	15	15%	46
NIG-58	110	2	DPL12	20	19%	69
WEIR	112	2	DPL11	26	25%	21
THY-32	111	1	DPL8	. 8	8%	27
TNF-H9G1	111	. 1	DPL8	9	9%	27
mAb61	111	1	DPL3	. 1	1%	29
LV1L1	98	1	DPL2	0	0%	54
НА	113	1	DPL3	14	13%	63
LA1L1	111	1	DPL2	3	3%	54
RHE	112	1	DPL1	17	16%	22
K1B12L	113	1	DPL8	17	16%	79
LOC	113	1	DPL2	15	14%	. 84
NIG-51	112	1	DPL2	12	11%	67
NEWM	104	1	DPL8	23	22%	10
MD3-4	106	3	DPL23	14	13%	4
COX	112	1	DPL2	13	12%	84
HiH10	106	3	DPL23	13	12%	3
VOR	112	.1	DPL2	16	15%	16
AL POL	113	1	DPL2	16	15%	57
CD4-74	111	1	DPL2	19	18%	27
AMYLOID MOL	102	3	DPL23	15	15%	30
OST577	108	3	Humlv318	10	10%	4
NIG-48	113	1	DPL3	42	40%	66
CARR	108	3	DPL23	18	17%	19
			66			

Table 2B: (continued)

Name <sup>1</sup>	aa²	Computed family <sup>3</sup>	Germline gene <sup>4</sup>	Diff. to germline <sup>s</sup>	% diff. to germline <sup>6</sup>	Reference'
mAb60	108	3	DPL23	14	13%	29
NIG-68	99	3	DPL23	25	26%	32
KERN	107	3	DPL23	26	25%	59
ANT	106	3	DPL23	17	16%	19
LEE	110	3	DPL23	18	17%	85
CLE ·	94	3	DPL23	17	17%	19
VL8	98	8	DPL21	0	0%	81
MOT	110	3	Humlv318	23	22%	38
GAR	108	. 3	DPL23	26	25%	33
32.B9	98	8	DPL21	5	5%	81
PUG	108	3	Humlv318	24	23%	19
T1	115	8	HUMLV801	52	50%	6
RF-TS7	96	. 7	DPL18	4	40/0	60
YM-1	116	8	HUMLV801	51	49%	75
K6H6	112	8	HUMLV801	20	19%	44
K5C7	112	8	HUMLV801	20	19%	44
K5B8	112	8	HUMLV801	20	19%	44
K5G5	112	8	HUMLV801	20	19%	44
K4B8	112	. 8	HUMLV801	19	18%	44
K6F5	112	8	HUMLV801	17	16%	44
HIL	108	3	DPL23	22	21%	47
KIR	109	3	DPL23	20	19%	19
CAP.	109	3	DPL23	19	18%	84
1B8	110	3	DPL23	22	21%	- 43
SHO	108	3	DPL23	19	18%	19
HAN	108	. 3	DPL23	20	19%	19
cML23	96	3	DPL23	3	3%	12
PR-SJ1	96	3	DPL23	7	7%	55
BAU	107	3	DPL23	9	9%	5
TEX	99	3	DPL23	8	8%	19
X(PET)	107	3	DPL23	9	9%	51
DOY	106	3	DPL23	9	9%	19
COT	106	3	DPL23	13	12%	19
Pag-1	111	3	Humlv318	5	5%	31

Table 2B: (continued)

Name <sup>1</sup>	aa²	Computed family <sup>3</sup>	Germline gene <sup>4</sup>	Diff. to germline <sup>s</sup>	% diff. to germline <sup>6</sup>	Reference'
DIS	107	3	Humlv318	2	2%	19
WIT	108	3	Humlv318	. 7	7%	19
I.RH	108	3	Humlv318	12	11%	19
S1-1	108	3	Humlv318	12	11%	52
DEL	108	3	Humlv318	14	13%	. 17
TYR	108	3	Humlv318	11	10%	19
J.RH	109	3	Humlv318	13 ·	12%	19
THO	112	2	DPL13	38	36%	26
LBV	113	1	DPL3	38	36%	2
WLT	112	1	DPL3	33	31%	14
SUT	112	2	DPL12	37	35%	65

Table 2C: rearranged human heavy chain sequences

Name <sup>1</sup>	aa²	Computed family <sup>3</sup>	Germline gene⁴	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference <sup>3</sup>
21/28	119	1	VH1-13-12	0	0,0%	31
8E10	123	1	VH1-13-12	0	0.0%	31
MUC1-1	118	1	VH1-13-6	4	4,1%	42
gF1	98	1	VH1-13-12	10	10,2%	75
VHGL 1.2	98	1	VH1-13-6	2	2,0%	26
HV1L1	98	1	VH1-13-6	0	0,0%	81
RF-TS7	104	1	VH1-13-6	3	3,1%	96
E55 1.A15	106	1	VH1-13-15	1	1,0%	26
HA1L1	126	· 1	VH1-13-6	7	7,1%	81
UC	123	1	VH1-13-6	5	5,1%	115
WIL2	123	1	VH1-13-6	6	6,1%	55
R3.5H5G	122	1	VH1-13-6	10	10,2%	70
N89P2	1,23	1	VH1-13-16	11	11,2%	· 77
mAb113	126	1	VH1-13-6	10	10,2%	71
LS2S3-3	125	1	VH1-12-7	5	5,1%	98
LS2S3-12a	125	1	VH1-12-7	· <b>5</b>	5,1%	98
LS2S3-5	125	1	VH1-12-7	5	5,1%	98
LS2S3-12e	125	. 1	VH1-12-7	5	5,1%	98
LS2S3-4	125	1	VH1-12-7	<b>5</b> .	5,1%	98
LS2S3-10	. 125	1	VH1-12-7	5	5,1%	98
LS2S3-12d	125	1	VH1-12-7	6	6,1%	98
LS2S3-8	125	1	VH1-12-7	5	5,1%	98
LS2	125	1	VH1-12-7	6	6,1%	113
LS4	105	1	VH1-12-7	6	6,1%	113
LS5	125	1	VH1-12-7	6	6,1%	113
LS1	125	1	VH1-12-7	6	6,1%	113
LS6	125	1	VH1-12-7	6	6,1%	113
LS8	125	· · 1	VH1-12-7	7	7.1%	113
THY-29	122	1	VH1-12-7	0	0,0%	42
1B9/F2	122	1	VH1-12-7	10	10,2%	21
51P1	122	1	VH1-12-1	0	0.0%	105
NEI	127	1	VH1-12-1	0	0.0%	55
AND	127	1	VH1-12-1	0	0.0%	55
L7	127	1	VH1-12-1	0	0,0%	54
L22	124	1	VH1-12-1	0	0,0%	54
L24	127	1	VH1-12-1	0	0,0%	54

Table 2C: (continued)

Name¹	aa²	Computed family <sup>3</sup>	Germline gene⁴	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference <sup>2</sup>
L26	116	1 .	VH1-12-1	0	0,0%	54
L33	119	1	VH1-12-1	0	0,0%	54
L34	117	1	VH1-12-1	. 0	0,0%	54
L36	118	1 .	VH1-12-1	0	0.0%	<b>54</b> .
L39	120	1	VH1-12-1	0	0,0%	54
L41 .	120	1	VH1-12-1	0	0,0%	54
L42	125	. 1	VH1-12-1	0	0,0%	54
VHGL 1.8	101	1	VH1-12-1	0	0,0%	26
783c	127	1	VH1-12-1	0	0,0%	22
X17115	127	1	VH1-12-1	0	0,0%	37
L25	124	1	VH1-12-1	0	0,0%	54
L17	120	1	VH1-12-1	1	1,0%	54
L30	127	1	VH1-12-1	1	1,0%	54
L37	120	1	VH1-12-1	· 1	1,0%	54
TNF-E7	116	1 .	VH1-12-1	2	2,0%	42
mAb111	122	1	VH1-12-1	7.	7,1%	71
III-2R	122	1	VH1-12-9	3	3,1%	70
KAS	121	1	VH1-12-1	7	7,1%	79
YES8c	122	1	VH1-12-1	8	8,2%	34
RF-TS1	123	1	VH1-12-1	8	8,2%	82
BOR'	121	1	VH1-12-8	. 7	7,1%	79
VHGL 1.9	101	1 .	VH1-12-1	8	8,2%	26
mAb410.30F305	117	1	VH1-12-9	5	5,1%	52
EV1-15	127	1	VH1-12-8	10	10,2%	78
mAb112	122	1	VH1-12-1	11	11,2%	71
EU	117	1	VH1-12-1	11	11,2%	28
H210	127	1	VH1-12-1	12	12,2%	66
TRANSGENE	104	1	VH1-12-1	0	0,0%	111
CLL2-1	93	1	VH1-12-1	0	0,0%	30
CLL10 13-3	97	1	VH1-12-1	0 .	0,0%	29
LS7	99	1	VH1-12-7	4	4,1%	. 113
ALL7-1	87	1 .	VH1-12-7	0	0,0%	30
CLL3-1	91	1	VH1-12-7	1	1,0%	30
ALL56-1	85	1	VH1-13-8	0	0,0%	30
ALL1-1	87	1	VH1-13-6	1	1,0%	30
ALL4-1	94	1	VH1-13-8	0	0,0%	30

Table 2C:

Name <sup>1</sup>	aa²	Computed family <sup>3</sup>	Germline gene⁴	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference <sup>7</sup>
ALL56 15-4	85	1	VH1-13-8	5	5,1%	29
CLL4-1	88	1	VH1-13-1	1	1,0%	. 30
Au92.1	98	1	VH1-12-5	0	0,0%	49
RF-TS3	120	1	VH1-12-5	1	1,0%	82
Au4.1	98	1	VH1-12-5	1	1,0%	49
HP1	121	1	VH1-13-6	13	13,3%	110
BLI	127	1	VH1-13-15	5	5,1%	72
No.13	127	·, 1	VH1-12-2	19	19,4%	76
TR1.23	122	1	VH1-13-2	23	23,5%	88
S1-1	125	1	VH1-12-2	18	18,4%	76
TR1.10	119	1	VH1-13-12	14	14,3%	88
E55 1.A2	102	1 .	VH1-13-15	3	3.1%	<b>26</b> .
SP2	119	1	VH1-13-6	15	15,3%	89
TNF-H9G1	111	1	VH1-13-18	2	2,0%	42
G3D10H	127	1	VH1-13-16	19	19,4%	127
TR1.9	118	1	VH1-13-12	14	14,3%	88
TR1.8	121	1	VH1-12-1	24	24,5%	88
LUNm01	127	1	VH1-13-6	22	22,4%	9
K1B12H	127	. 1	VH1-12-7	23	23,5%	127
L3B2	99	1	VH1-13-6	. 2	2,0%	46
ss2	100	1	VH1-13-6	2	2,0%	46
No.86	124	1	VH1-12-1	20	20,4%	76
TR1.6	124	1	VH1-12-1	19	19,4%	88
ss7	99	1	VH1-12-7	- 3	3,1%	46
s5B7	102	1	VH1-12-1	0,	0,0%	46
s6A3	97	1	VH1-12-1	0	0,0%	46
ss6	99	1	VH1-12-1	0	0,0%	46
L2H7	103	1	VH1-13-12	0	0,0%	46
s6BG8	93	1	VH1-13-12	0	0.0%	46
s6C9	107	1	VH1-13-12	0	0,0%	46
HIV-b4	124	1	VH1-13-12	21	21,4%	12
HIV-b12	124	1	VH1-13-12	21	21,4%	12
L3G5	98	1	VH1-13-6	1	1,0%	46
22	115	1	VH1-13-6	11	11,2%	118
L2A12	99	1	VH1-13-15	3	3,1%	46
PHOX15	124	. 1	VH1-12-7	20	20,4%	73
	•		71			

Name <sup>1</sup>	aa²	Computed family <sup>3</sup>	Germline gene <sup>4</sup>	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference <sup>7</sup>
LUNm03	127	1	VH1-1X-1	18	18,4%	9
CEA4-8A	129	1	VH1-12-7	1	1,0%	42
M60	121	<b>2</b> .	VH2-31-3	3	3,0%	103
HiH10	127	2	VH2-31-5	. 9	9,0%	4
COR	119	2	VH2-31-2	11 .	11,0%	91
2-115-19	124	2 .	VH2-31-11	8	8,1%	124
OU	125	2	VH2-31-14	20	25,6%	92
HE	120	2	VH2-31-13	19	19,0%	27
CLL33 40-1	78	2	VH2-31-5	2	2,0%	29
E55 3.9	88	3	VH3-11-5	7	7,2%	26
MTFC3	125	3	VH3-14-4	21	21,0%	131
MTFC11	125	3	VH3-14-4	21	21,0%	131
MTFJ1	114	3	VH3-14-4	21	21,0%	131
MTFJ2	114	3	VH3-14-4	21	21,0%	131
MTFUJ4	100	3	VH3-14-4	21	21,0%	131
MTFUJ5	100	3	VH3-14-4	21	21,0%	131
MTFUJ2	100	3 -	VH3-14-4	22	22,0%	131
MTFC8	125	3.	VH3-14-4	23	23,0%	131
TD e Vq	113	3	VH3-14-4	0	0,0%	16
rMTF	. 114	3	VH3-14-4	5	5,0%	131
MTFUJ6	100	3	VH3-14-4	· 10	10,0%	131
RF-KES	107	3	· VH3-14-4	. 9	9,0%	85
N51P8	126	3	VH3-14-1	9	9.0%	77
TEI	119	3	VH3-13-8	21	21,4%	20
33.H11	115	3	VH3-13-19	- <b>10</b>	10,2%	129
SB1/D8	101	3	VH3-1X-8	14	14,0%	2
38P1	119	3	VH3-11-3	0	0,0%	104
BRO'IGM	119	3	VH3-11-3	13	13,4%	19
NIE	119	3	VH3-13-7	15	15,3%	87
3D6	126	3	VH3-13-26	5	5,1%	35
ZM1-1	112	3	VH3-11-3	8	8,2%	5
E55 3.15	110	3	VH3-13-26	0	0,0%	26
gF9	108	3	VH3-13-8	15	15,3%	75
THY-32	120	3	VH3-13-26	3	3,1%	42
RF-KL5	100.	3	VH3-13-26	5	5,1%	96
OST577	122	3	VH3-13-13	6	6,1%	5

Name <sup>1</sup>	aa²	Computed family <sup>3</sup>	Germline gene⁴	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference
ВО	- 113	3	VH3-13-19	15	15,3%	10
Π125	121	3	VH3-13-10	15	15,3%	64
2-115-58	127	3	VH3-13-10	11	11,2%	. 124
KOL	126	3	VH3-13-14	16	16,3%	102
mAb60	118	3	VH3-13-17	14	14,3%	45
RF-AN	106	. 3	VH3-13-26	. 8	8,2%	85
BUT	115	3	VH3-11-6	13	13,4%	119
KOL-based CAMPATH						
9	118	3	VH3-13-13	16	16,3%	41
B1	119	3	VH3-13-19	13	13,3%	53
N98P1	127	3	VH3-13-1	13	13,3%	77
Π117	107	3	VH3-13-10	12	12,2%	64
WEA	114	3	VH3-13-12	15	15,3%	40
HIL	120	3	VH3-13-14	14	14,3%	23
s5A10	97	3	VH3-13-14	0	0,0%	46
s5D11	98	3	VH3-13-7	0 .	0,0%	46
s6C8	100	3	VH3-13-7	0	0,0%	46
s6H12	98	3	VH3-13-7	0	0,0%	46
VH10.7	119	3	VH3-13-14	16	16,3%	128
HIV-loop2	126	3	VH3-13-7	16	16,3%	12
HIV-loop35	126	3	VH3-13-7	16	16,3%	12
TRO	122	<b>3</b> .	VH3-13-1	13	13,3%	61
SA-4B	123	3	VH3-13-1	15	15,3%	125
L2B5	98	3	VH3-13-13	0	0,0%	46
s6E11	95	3	VH3-13-13	0	0.0%	46
s6H7	100	3	VH3-13-13	0	0,0%	46
ss1	102	3	VH3-13-13	0	0,0%	46
ss8	94	3	VH3-13-13	0	0,0%	46
DOB	120	3	VH3-13-26	21	21,4%	116
THY-33	115	3	VH3-13-15	20	20,4%	42
NOV	118	3	VH3-13-19	14	14,3%	38
rsv13H	120	3	VH3-13-24	20	20,4%	11
L3G11	98	3	VH3-13-20	. 2	2,0%	. 46
L2E8	99	3	VH3-13-19	0	0,0%	46
L2D10	101	3	VH3-13-10	1	1,0%	46
L2E7	98	3	VH3-13-10	1	1,0%	46

Table 2C: (continued)

Name¹	aa²	Computed family <sup>3</sup>	Germline gene⁴	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference'
L3A10	100	3	VH3-13-24	0	0,0%	46
L2E5	97	3	VH3-13-2	1	1,0%	46
BUR	119	3	VH3-13-7	21	21,4%	67
s4D5	107	3	VH3-11-3	1	1,0%	46
19	116	3	VH3-13-16	4	4,1%	118
s5D4	99	3	VH3-13-1	0	0,0%	46
s6A8	100	3	VH3-13-1	0	0,0%	46
HIV-loop13	123	3	VH3-13-12	17	17,3%	12
TR1.32	112	3	VH3-11-8	18	18,6%	88
L2B10	97	3	VH3-11-3	1	1,0%	46
ŢR1.5	114	3	VH3-11-8	21	21,6%	88
s6H9	101	3	VH3-13-25	0	0.0%	46
8 .	112	3	VH3-13-1	6	6,1%	118
23	115	3	VH3-13-1	6	6,1%	118
7	115	3	VH3-13-1	4	4,1%	118
TR1.3	120	3	VH3-11-8	20	20,6%	88
18/2	125	3	VH3-13-10	0	0,0%	32
18/9	125	3	VH3-13-10	0	0,0%	. 31
30P1	. 119	3	VH3-13-10	0	0,0%	106
HF2-1/17	125	3	VH3-13-10	0	0,0%	8
A77	109	3	VH3-13-10	0	0,0%	44
B19.7	108	3 .	VH3-13-10	. 0	0,0%	44
M43	119	3	VH3-13-10	0	0,0%	103
1/17	125	3	VH3-13-10	0	0,0%	31
18/17	125	3	VH3-13-10	0	0,0%	<b>31</b> .
E54 3.4	109	3	VH3-13-10	0	0,0%	26
LAMBDA-VH26	98	3	VH3-13-10	1	1,0%	95
E54 3.8	111	3	VH3-13-10	1	1,0%	26
GL16	106	3	VH3-13-10	1	1,0%	44
4G12	125	3	VH3-13-10	1	1,0%	56
A73	106	3	VH3-13-10	2	2,0%	44
AL1.3	111	3	VH3-13-10	3	3,1%	117
3.A290	118	3	VH3-13-10	2	2,0%	108
Ab18	127	3	VH3-13-8	2	2,0%	100
E54 3.3	105	3	VH3-13-10	3	3,1%	26
35G6	121	3	VH3-13-10	3	3,1%	57

Name <sup>1</sup>	aa²	Computed family <sup>3</sup>	Germline gene⁴	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference'
A95	107	3	VH3-13-10	5	5,1%	44
Ab25	128	3	VH3-13-10	5	5,1%	100
N87	126	3	VH3-13-10	4	4,1%	77
ED8.4	99	3	VH3-13-10	6	6,1%	2
RF-KL1	122	3	VH3-13-10	6	6,1%	82
AL1.1	112	3	VH3-13-10	2	2,0%	117
AL3.11	102	3	VH3-13-10	1	1,0%	117
32.B9	127	3	VH3-13-8	6	6,1%	129—
TK1	109	3	VH3-13-10	2	2,0%	117
POP	123	3	VH3-13-10	8	8,2%	115
9F2H	127	3	VH3-13-10	9	9,2%	127
VD	115	3	VH3-13-10	9	9,2%	10
Vh38Ci.10	121	3	VH3-13-10	8	8,2%	74 ·
Vh38Cl.9	121	3	VH3-13-10	8	8,2%	74
Vh38Cl.8	121	3	VH3-13-10	8	8,2%	74
63P1	120	3	VH3-11-8	0	0,0%	104
60P2	117	3	VH3-11-8	0	0.0%	104
AL3.5	90	3	VH3-13-10	<b>' 2</b>	2,0%	117
GF4/1.1	123	3	VH3-13-10	10	10,2%	39
Ab21	126	3	VH3-13-10	12	12,2%	100
TD d Vp	118	3	VH3-13-17	2	2,0%	16
Vh38Cl.4	119	3	VH3-13-10	8	8,2%	74
Vh38Cl.5	119	3	VH3-13-10	8	8,2%	74
AL3.4	104	3	VH3-13-10	1	1,0%	117
FOG1-A3	115	3	VH3-13-19	2	2,0%	42.
HA3D1	117	3	VH3-13-21	1	1,0%	81
E54 3.2	112	3	VH3-13-24	0	0,0%	26
mAb52	128	3	VH3-13-12	2	2,0%	51
mAb53	128	. 3	VH3-13-12	2	2,0%	51
mAb56	128	3	VH3-13-12	2	2,0%	51
mAb57	128	3	VH3-13-12	2	2,0%	51
mAb58	128	3	VH3-13-12	2	2,0%	51
mAb59	128	3	VH3-13-12	2	2,0%	51
mAb105	128	3	VH3-13-12	2	2,0%	51
mAb107	128	3	VH3-13-12	2	2,0%	51
E55 3.14 .	110	3	VH3-13-19	0	0,0%	26

Table 2C:

Name <sup>1</sup>	aa²	Computed family <sup>3</sup>	Germline gene⁴	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference <sup>7</sup>
F13-28	106	3	VH3-13-19	1	1,0%	. 94
mAb55	127	3	VH3-13-18	4	4,1%	51
YSE <sup>-</sup>	117	3	VH3-13-24	6	6,1%	72
E55 3.23	106	3	VH3-13-19	2	2,0%	26
RF-TS5	101	3	VH3-13-1	3	3,1%	85
N42P5	124	3	VH3-13-2	7	7,1%	77
FOG1-H6	110	- 3	VH3-13-16	7	7,1%	42
0-81	115	3	VH3-13-19	11	11,2%	47
HIV-s8	122	3	VH3-13-12	11	11,2%	. 12
mAb114	125	3	VH3-13-19	12	12,2%	71
33.F12	116	3	VH3-13-2	4	4,1%	129
4B4	119	3	VH3-1X-3	0	0,0%	101
M26	123	3	VH3-1X-3	0	0,0%	103
VHGL 3.1	100	3	VH3-1X-3	0 .	0,0%	<b>26</b> ·
E55 3.13	113	3	VH3-1X-3	1	1,0%	26
SB5/D6	101	3	VH3-1X-6	3	3,0%	2
RAY4	101	3	VH3-1X-6	3	3,0%	2
82-D V-D	106	3	VH3-1X-3	5	5,0%	112
MAL	129	3	VH3-1X-3	5	5,0%	72
LOC	123	3	VH3-1X-6	5	5,0%	72
LSF2	101	3	VH3-1X-6	11	11,0%	2
HIB RC3	100	3	· VH3-1X-6	11	11,0%	1
56P1	119	3	VH3-13-7	0	0,0%	104
M72	122	3	VH3-13-7	0	0,0%	103
M74	121	3	VH3-13-7	0	0,0%	103
E54 3.5	105	3	VH3-13-7	0	0,0%	26
2E7	123	3	VH3-13-7	0	0.0%	63
2P1	117	3	VH3-13-7	0	0,0%	104
RF-SJ2	127	3	VH3-13-7	1	1,0%	83
PR-TS1	114	3	VH3-13-7	1	1,0%	85
KIM46H	127	3	VH3-13-13	0	0,0%	18
E55 3.6	108	3	VH3-13-7	2	2,0%	26
E55 3.10	107	3	VH3-13-13	1	1,0%	26
3.B6	114	3	VH3-13-13	1	1,0%	108
E54 3.6	110	3	VH3-13-13	1	1,0%	26
FL2-2	114	3	VH3-13-13	1	1,0%	80

Table 2C:

Name <sup>1</sup>	aa²	Computed family <sup>3</sup>	Germline gene⁴	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference <sup>2</sup>
RF-SJ3	112	. 3	VH3-13-7	2	2,0%	85
E55 3.5	105	3	VH3-13-14	1	1,0%	26
BSA3	121.	3	VH3-13-13	1	1,0%	73
HMST-1	119	3	VH3-13-7	<b>3</b> .	3,1%	130
RF-TS2	126	3	VH3-13-13	4	4,1%	82
E55 3.12	109	3	VH3-13-15	0	0,0%	26
19.E7	126	3	VH3-13-14	3	3,1%	129
11-50	119	3	VH3-13-13	6	6,1%	130
E29.1	120	3	VH3-13-15	2	2,0%	25
E55 3.16	108	3	VH3-13-7	6	6,1%	. 26
TNF-E1	117	3	VH3-13-7	7	7,1%	42
RF-SJ1	127	3	VH3-13-13	6	6,1%	83
FOG1-A4	116	3	VH3-13-7	8	8,2%	42
TNF-A1	117	3	VH3-13-15	4	4,1%	42
PR-SJ2	107	3	VH3-13-14	8	8,2%	85
HN.14	124	3	VH3-13-13	10	10,2%	33
CAM'	121	3	VH3-13-7	12	12,2%	65
HIV-B8	125	3	VH3-13-7	9	9,2%	12
HIV-b27	125	3	VH3-13-7	9	9,2%	12
HIV-b8	125	<b>3</b>	VH3-13-7	9	9,2%	12
HIV-s4	125	3	VH3-13-7	9	9,2%	12
HIV-B26	125	3	VH3-13-7	9	9,2%	12
HIV-B35	125	3	VH3-13-7	10	10,2%	12
HIV-b18	125	3	VH3-13-7	10	10,2%	12
HIV-b22	125	3	VH3-13-7	11	11,2%	.12
HIV-b13	125	3	VH3-13-7	12	12,2%	12
333	117	3	VH3-14-4	24	24,0%	24
1H1	120	3	VH3-14-4	24	24,0%	24
1B11	120	3	VH3-14-4	23	23,0%	24
CLL30 2-3	86	3	VH3-13-19	1	1,0%	29
GA	110	3	VH3-13-7	19	19,4%	36
JeB	99	3	VH3-13-14	3	3,1%	7
GAL	110	3	VH3-13-19	10	10,2%	126
К6Н6	119		VH3-1X-6	18	18,0%	60
K4B8	119	3	VH3-1X-6	18	18,0%	60
K5B8	119	3	VH3-1X-6	18	18,0%	60

Table 2C: (continued)

Name <sup>1</sup>	aa²	Computed family <sup>3</sup>	Germline gene <sup>4</sup>	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference'
K5C7	119	3	VH3-1X-6	19	19,0%	60
K5G5	119	3	VH3-1X-6	19	19,0%	60
K6F5	119	3	VH3-1X-6	19	19,0%	60
AL3.16	98	3	VH3-13-10	1	1,0%	117
N86P2	98	3	VH3-13-10	3	3,1%	77
N54P6	95	3	VH3-13-16	7	7,1%	77
LAMBDA HT112-1	126	4	VH4-11-2	0	0,0%	. 3
HY18	121	4	VH4-11-2	0	0.0%	43—
mAb63	126	4	VH4-11-2	0	0,0%	45
FS-3	105	4	VH4-11-2	0	0,0%	86
FS-5	111	4	VH4-11-2	0	0,0%	86
FS-7	107	4	VH4-11-2	0	0,0%	86
FS-8	110	4	VH4-11-2	0	0,0%	86
PR-TS2	105	4	VH4-11-2	0	0,0%	85
RF-TMC	102	4	VH4-11-2	0	0,0%	85
mAb216	122	4	VH4-11-2	1	1,0%	15
mAb410.7.F91	122	4	VH4-11-2	1	1,0%	52
mAbA6H4C5	124	4	VH4-11-2	1	1,0%	15
Ab44	127	4 .	VH4-11-2	2	2,1%	100
6H-3C4	124	4 .	VH4-11-2	3	3,1%	59
FS-6	108	4	VH4-11-2	. 6	6,2%	86
FS-2	114	4 .	VH4-11-2	6	6,2%	84
HIG1	126	4	VH4-11-2	7	7,2%	62
FS-4	105	4	VH4-11-2	8	8,2%	86
SA-4A	123	4	VH4-11-2	9	9,3%	125
LES-C	119	4	VH4-11-2	10	10,3%	99
DI	78	4	VH4-11-9	16	16,5%	58
Ab26	126	4	VH4-31-4	8	8,1%	100
TS2	124	4	VH4-31-12	15	15,2%	110
265-695	115	4	VH4-11-7	16	16,5%	5
WAH	129	4	VH4-31-13	19	19,2%	93
268-D	122	4	VH4-11-8	22	22,7%	6
58P2	118	4	VH4-11-8	. 0	0,0%	104
mAb67	128	4	VH4-21-4	1	1,0%	45
4.L39	115	4	VH4-11-8	. 2	2,1%	108
mF7	111	4.	VH4-31-13	3	3,0%	75

Table 2C:

Name¹	aa²	Computed family <sup>3</sup>	Germline gene⁴	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference <sup>7</sup>
33.C9	122	4	VH4-21-5	7	7,1%	129
Pag-1	124	4	VH4-11-16	5	5,2%	50
<b>B3</b>	123	4	VH4-21-3	8	8,2%	53
IC4	120	4	VH4-11-8	6	6,2%	70
C6B2	127	4	VH4-31-12	4	4,0%	48
N78	118	4	VH4-11-9	11	11,3%	77
B2	109	4	VH4-11-8	12	12,4%	53
WRD2	123	4	VH4-11-12	6	6,2%	90
mAb426.4.2F20	126	4	VH4-11-8	· 2	2,1%	52
E54 4.58	115	4	VH4-11-8	1	1,0%	26
WRD6	123	4	VH4-11-12	10	10,3%	90
mAb426.12.3F1.4	122	4	VH4-11-9	· <b>4</b>	4,1%	52
E54 4.2	108	4	VH4-21-6	2	2,0%	26
WIL	127	4	VH4-31-13	0 .	0,0%	90
COF	126	4	VH4-31-13	0	0,0%	90
LAR	122	4	VH4-31-13	2	2,0%	90
WAT	125	4	VH4-31-13	4	4,0%	90
mAb61	123	4	VH4-31-13	5	5,1%	45
WAG	127	4	VH4-31-4	0	0,0%	90
RF-SJ4	108	4	VH4-31-12	2	2,0%	85
E54 4.4	110	4	VH4-11-7	0	0,0%	26
E55 4.A1	108	4	VH4-11-7	0	0,0%	26
PR-SJ1	103	4	VH4-11-7	1	1,0%	85
E54 4.23	111	4	VH4-11-7	1	1,0%	26
CLL7 7-2	97	4	VH4-11-12	0	0,0%	29
37P1	95	4	VH4-11-12	0	0,0%	104
ALL52 30-2	91	4	VH4-31-12	4	4,0%	29
EBV-21	98	5	VH5-12-1	0	0,0%	13
CB-4	98	5	VH5-12-1	0	0,0%	13
CLL-12	98	5	VH5-12-1	0	0,0%	13
L3-4	98	5	VH5-12-1	0	0,0%	13
CLL11	98	5	VH5-12-1	0	0,0%	17
CORD3	98	5	VH5-12-1	0	0,0%	17
CORD4	98	5	VH5-12-1	0	0,0%	17
CORD8	98	5	VH5-12-1	0	0,0%	17
CORD9	98	5	VH5-12-1	0	0,0%	17

Table 2C:

Name¹	aa²	Computed family <sup>3</sup>	Germline gene⁴	Diff. to germline <sup>s</sup>	% diff. to germline <sup>6</sup>	Reference <sup>2</sup>
CD+1	98	5 .	VH5-12-1	0	0,0%	17
CD+3	98	5	VH5-12-1	0	0,0%	- 17
CD+4	98	5	VH5-12-1	0	0,0%	17
CD-1	98	5	VH5-12-1	0	0,0%	17
CD-5	98	5	VH5-12-1	Ö	0,0%	17
VERG14	98	5	VH5-12-1	0	0,0%	17
PBL1	98	5	VH5-12-1	0	0,0%	17
PBL10	98	5	VH5-12-1	0	0,0%	17
STRAb SA-1A	127	5	VH5-12-1	0	0,0%	125
DOB'	122	5	VH5-12-1	0	0,0%	97
VERG5	98	5	VH5-12-1	0	0,0%	17
PBL2	98	5	VH5-12-1	1	1,0%	17
Tu16	119	5	VH5-12-1	1 -	1,0%	49
PBL12	98	5	VH5-12-1	1	1,0%	17
CD+2	98	5	VH5-12-1	1	1,0%	17
CORD10	98	. 5	VH5-12-1	1	1,0%	17
PBL9	98	. 5	VH5-12-1	1	1,0%	17
CORD2	98	5	VH5-12-1	2	2,0%	17
PBL6	98	5	VH5-12-1	2	2,0%	17
CORD5	98	5	VH5-12-1	2	2,0%	17
CD-2	98	5	VH5-12-1	. 2	2,0%	17
CORD1	98	5	VH5-12-1	2 .	2,0%	. 17
CD-3	98	5	VH5-12-1	3	3,1%	17
VERG4	98	5	VH5-12-1	3	3,1%	<b>17</b> .
PBL13	98	.5	VH5-12-1	3	3,1%	17
PBL7	98	5	VH5-12-1	3	3,1%	17
HAN	119	5	VH5-12-1	3	3,1%	97
VERG3	98	· 5	VH5-12-1	3	3,1%	17
PBL3	98	5	VH5-12-1	<b>3</b> ·	3,1%	17
VERG7	98	5	VH5-12-1	3	3,1%	17
PBL5	94	5	VH5-12-1	0	0,0%	17
CD-4	98	5	VH5-12-1	4	4,1%	17
CLL10	98	5	VH5-12-1	4	4,1%	17
PBL11	98	5	VH5-12-1	4	4,1%	17
CORD6	98	5	VH5-12-1	. 4	4,1%	17
VERG2	98	5	VH5-12-1	5	5,1%	17

Table 2C: (continued)

Name <sup>1</sup>	aa²	Computed family <sup>3</sup>	Germline gene⁴	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference <sup>7</sup>
83P2	119	5	VH5-12-1	0	0,0%	103
VERG9	98	. 5	VH5-12-1	6	6,1%	17
CLL'6	98	5	VH5-12-1	6	6,1%	17
PBL8	98	5	VH5-12-1	7	7,1%	17
Ab2022	120	5	VH5-12-1	3	3,1%	100
CAV	127	5	VH5-12-4	0	0,0%	97
HOW'	120	5	VH5-12-4	0	0,0%	97
PET	127	5	VH5-12-4	. 0	0,0%	97
ANG	121	5 -	VH5-12-4	0	0,0%	97
KER	121	5	VH5-12-4	0	0,0%	97
5.M13	118	5	VH5-12-4	0	0,0%	107
Au2.1	118	<b>5</b> ·	VH5-12-4	1	1,0%	49
WS1	126	5	VH5-12-1	9	9,2%	110
TD Vn	98	5	VH5-12-4	1	1,0%	16
TEL13	116	5	VH5-12-1	9	9,2%	73
E55 5.237	112	5	VH5-12-4	2	2,0%	26
VERG1	98	5	VH5-12-1	10	10,2%	17
CD4-74	117	5	VH5-12-1	10	10,2%	42
257-D	125	5	VH5-12-1	11	11,2%	6
CLL4	98	5	VH5-12-1	11	11,2%	17
CLL8	98	5	VH5-12-1	11	11,2%	17
Ab2	124	5	VH5-12-1	12	12,2%	120
Vh383ex	98	5	VH5-12-1	12	12,2%	120
CLL3	98	5	VH5-12-2	11	11,2%	17
Au59.1	122	5	VH5-12-1	12	12,2%	49
TEL16	117	5	VH5-12-1	12	12,2%	73
M61	104	5	VH5-12-1	0	0,0%	103
Tu0	99	5	VH5-12-1	5	5,1%	49
P2-51	122	5	VH5-12-1	13	13,3%	121
P2-54	122	5	VH5-12-1	11	11,2%	121
P1-56	119	5	VH5-12-1	9	9,2%	121
P2-53	122	5	VH5-12-1	10	10,2%	121
P1-51	123	5	VH5-12-1	19	19,4%	121
P1-54	123	5	VH5-12-1	3	3,1%	121
P3-69	127	5	VH5-12-1	4	4,1%	121
P3-9	119	5	VH5-12-1	4	4,1%	121

Table 2C: (continued)

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Name¹	aa²	Computed family <sup>3</sup>	Germline gene⁴	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference
1-185-37	125	5 .	VH5-12-4	0	0,0%	124
1-187-29	. 125	5	VH5-12-4	0	0,0%	124
Pi-58	128	5	VH5-12-4	10	10,2%	121
P2-57	118	5	VH5-12-4	3	3,1%	121
P2-55	123	5	VH5-12-1	5	5,1%	121
P2-56	123	5	VH5-12-1	20	20,4%	121
P2-52	122	5	VH5-12-1	11	11,2%	121
P3-60	122	5	VH5-12-1	8	8,2%	121
P1-57	123	5	VH5-12-1	4	4,1%	121
P1-55	122	5	VH5-12-1	14	14,3%	121
MD3-4	128	5	VH5-12-4	12	12,2%	5
P1-52	121	5	VH5-12-1	11	11,2%	121
CLL5	98	5	VH5-12-1	13	13,3%	17
CLL7	98	5	VH5-12-1	14	14,3%	17
L2F10	100	5	VH5-12-1	1	1,0%	46
L3B6	98	5	VH5-12-1	1	1,0%	46
VH6.A12	119	6	VH6-35-1	13	12,9%	122
s5A9	102	6	VH6-35-1	1	1,0%	. 46
s6G4	99	6	VH6-35-1	1	1,0%	46
ss3	99	6	VH6-35-1	1 -	1,0%	46
6-1G1	101	6	VH6-35-1	0	0,0%	14
F19L16	107	6 ·	VH6-35-1	0	0,0%	68
L16	120	6	VH6-35-1	0	0,0%	69
M71	121	6	VH6-35-1	0	0,0%	103
ML1	120	6	VH6-35-1	0	0,0%	69
F19ML1	107	6	VH6-35-1	0	0,0%	68
15P1	127	6	VH6-35-1	0	0,0%	104
VH6.N1	121	. 6	VH6-35-1	0 .	0,0%	122
VH6.N11	123	6	VH6-35-1	0	0,0%	122
VH6.N12	123	6	VH6-35-1	0	0,0%	122
VH6.N2	125	6	VH6-35-1	0	0,0%	122
VH6.N5	125	6	VH6-35-1	0	0.0%	122
VH6.N6	127	6	VH6-35-1	0	0,0%	122
VH6.N7	126	6	VH6-35-1	0	0,0%	122
VH6.N8	123	6	VH6-35-1	0	0,0%	122
VH6.N9	123	6	VH6-35-1	0	0,0%	122

Table 2C: (continued)

18010 20.	(continued)	· .				
Name <sup>1</sup>	aa²	Computed family <sup>3</sup>	Germline gene <sup>4</sup>	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference'
VH6.N10	123	6	VH6-35-1	0	0,0%	122
VH6.A3	123	6	VH6-35-1	0	0,0%	122
VH6.A1	124	6	VH6-35-1	0	0,0%	122
VH6.A4	120	6	VH6-35-1	0	0,0%	122
E55 6.16	116	6	VH6-35-1	0	0,0%	26
E55 6.17	. 120	6	VH6-35-1	0	0,0%	26
E55 6.6	120	6	VH6-35-1	0	0,0%	26
VHGL 6.3	102	6	VH6-35-1	0	0,0%	26
CB-201	118	6	VH6-35-1	0	0,0%	109
VH6.N4	122	6	VH6-35-1	0	0,0%	122
E54 6.4	109	6	VH6-35-1	1	1,0%	26
VH6.A6	126	6	VH6-35-1	1	1,0%	122
E55 6.14	120	6	VH6-35-1	1 .	1,0%	26
E54 6.6	107	6	VH6-35-1	1	1,0%	26
E55 6.10	112	6	VH6-35-1	1	1,0%	26
E54 6.1	107	6	VH6-35-1	2	2.0%	26
E55 6.13	120	6	VH6-35-1	2	2,0%	26
E55 6.3	120	6	VH6-35-1	2	2,0%	26
E55 6.7	116	6	VH6-35-1	2	2,0%	26
E55 6.2	120	6	VH6-35-1	2	2,0%	26
E55 6.X	. 111	6	VH6-35-1	2 .	2,0%	26
E55 6.11	111,	6	VH6-35-1	3	3,0%	26
VH6.A11	118	6	VH6-35-1	3	3,0%	122
A10	107	6	VH6-35-1	3	3,0%	68
E55 6.1	120	6	VH6-35-1	4	4,0%	26
FK-001	124	6	VH6-35-1	4	4,0%	65
VH6.A5	121	6	VH6-35-1	.4	4,0%	122
VH6.A7	123	6	VH6-35-1	4	4,0%	122
HBp2	119	6	VH6-35-1	4	4,0%	4
Au46.2	123	6	VH6-35-1	5	5,0%	49
A431	106	6	VH6-35-1	5	5,0%	68
VH6.A2	120	6	VH6-35-1	5	5,0%	122
VH6.A9	125	6	VH6-35-1	. 8	7,9%	122
VH6.A8	118	6	VH6-35-1	10	9,9%	122
VH6-FF3	118	6	VH6-35-1	2	2,0%	123
VH6.A10	126	6	VH6-35-1	12	11,9%	122

Name <sup>1</sup>	aa²	Computed family <sup>3</sup>	Germline gene⁴	Diff. to germline <sup>5</sup>	% diff. to germline <sup>6</sup>	Reference <sup>7</sup>
VH6-EB10	117	6	VH6-35-1	3	3,0%	123
VH6-E6	119	<b>6</b> .	VH6-35-1	· 6	5,9%	123
VH6-FE2	121	6	VH6-35-1	6	5,9%	123
VH6-EE6	116	6	VH6-35-1	6	5,9%	123
VH6-FD10	118	6	VH6-35-1	6	5,9%	123
VH6-EX8	113	6	VH6-35-1	6	5,9%	123
VH6-FG9	121	6	VH6-35-1	8	7,9%	123
VH6-E5	116	6	VH6-35-1	9	8,9%	123
VH6-EC8	122	6	VH6-35-1	9	8,9%	123
VH6-E10	120	6	VH6-35-1	10	9,9%	123
VH6-FF11	122	6	VH6-35-1	11	10,9%	123
VH6-FD2	115	6	VH6-35-1	11	10,9%	123
CLL10 17-2	88	6	VH6-35-1	4	4,0%	29
VH6-BB11	94	6	VH6-35-1	4	4,0%	123
VH6-B41	93	. 6	VH6-35-1	7	6,9%	123
JU17	102	6	VH6-35-1	3	3,0%	114
VH6-BD9	96	. 6	VH6-35-1	11	10,9%	123
VH6-BB9	94	6	VH6-35-1	12	11,9%	123

Table 3A: assignment of rearranged V kappa sequences to their germline counterparts

Family <sup>1</sup>	Name	Rearranged <sup>2</sup>	Sum
1	VkI-I	28	<del> </del>
1	Vk1-2	0	
1	Vk1-3	1	
1	Vk1-4	0	
1	Vk1-5	7	•
1	Vk 1-6	0	
1	Vk1-7	0	•
1	Vk1-8	2	
1	Vk1-9	9	
1	Vk1-10	0	
1	Vk1-11	1	
1	Vk1-12	7	
. 1	Vk1-13	1	
1	Vk!-14	7	
1	Vk1-15	2	
!	Vk1-16	2	
j	Vk1-17	16	
1	Vk1-18	1	
1	Vk1-19	33	
1	Vk1-20	1	•
1	Vk1-21	1	
1 ·	Vk1-22	0	
1	Vk1-23	0	119 entries
2	Vk2-1	0	·
2	Vk2-2	1	
2	Vk2-3	0	
2	Vk2-4	0	
2	Vk2-5	0	•
2	Vk2-6	.16	
2	Vk2-7	0	
2	Vk2-8	0	
2	Vk2-9	1	
2	Vk2-10	0	
2	Vk2-11	7	
2	Vk2-12	0	25 entries
3	Vk3-I	ſ	

Table 3A:

Family 1	Name	Rearranged <sup>2</sup>	Sum
3	Vk3-3	35	
3	Vk3-4	115	
3	Vk3-5	0	
. 3	Vk3-6	0	
. 3	Vk3-7	1	
3	Vk3-8	40	192 entries
4	Vk4-1	33	33 entries
5	Vk5-1	1	1 entry
6	Vk6-1	0	
6	Vk6-2	0	0 entries
7	Vk7-1	0	0 entries

Table 3B: assignment of rearranged V lambda sequences to their germline counterparts

Family <sup>1</sup>	Name	Rearranged <sup>2</sup>	Sum
1	DPL1	1.	
1	DPL2	14	•
1	DPL3	6	
1	DPL4	· <b>1</b>	•
1	HUMLV117	4	
1	DPL5	13	
1 .	DPL6	0	
1	DPL7	, <b>O</b>	
1	DPL8	3	
1	DPL9	0	42 entries
2	DPL10	5	······································
2	VLAMBDA 2.1	0	
2	DPL11	23	
2 .	DPL12	15	
2	DPL13	0	
2	DPL14	0	43 entries
3	DPL16	10	
3	DPL23	19	
3 .	Humlv318	9	38 entries
7	DPL18	1	
7	DPL19	0	1 entries
8	DPL21	2	
8	HUMLV801	6	8 entries
9	DPL22	0.	0 entries
unassigned	DPL24	0	0 entries
10	gVLX-4.4	0	0 entries

Table 3C: assignment of rearranged V heavy chain sequences to their germline counterparts

Family <sup>1</sup>	Name	Rearranged <sup>2</sup>	Sum
1	VH1-12-1	38	<u>-</u>
1	VH1-12-8	2	
1	VH1-12-2	2	
1	VH1-12-9	2	
1	VH1-12-3	0	
1	VH1-12-4	0 .	
1	. VH1-12-5	3	
1	VH1-12-6	0	
1	VH1-12-7	23	
· 1	VH1-13-1	1	
1.	VH1-13-2	1	
1	VH1-13-3	0	
1	VH1-13-4	0	
1	VH1-13-5	0	
1	VH1-13-6	17	
1	VH1-13-7	0	
1	VH1-13-8	3	
1	VH1-13-9	0	
1	VH1-13-10	0	
1 ,	VH1-13-11	0	
1	VH1-13-12	10	•
1 .	VH1-13-13	0	
1	VH1-13-14	0	
1	VH1-13-15	4	
1	VH1-13-16	2	
1	VH1-13-17	0	
1	VH1-13-18	1	
1	VH1-13-19	0	
1	VH1-1X-1	1	110 entries
2	VH2-21-1	0	
2	VH2-31-1	0	
2	VH2-31-2	. 1	
2	VH2-31-3	1.	
2	VH2-31-4	0	
2	VH2-31-5	2	
2	VH2-31-6	0	
2	VH2-31-7	. 0	

Table 3C: (continued)

	Name	Rearranged <sup>2</sup>	Sum
2	VH2-31-14	1	
2	VH2-31-8	0	
2	VH2-31-9	0	
2	VH2-31-10	0	
2	VH2-31-11	1	
2	VH2-31-12	0	
2	VH2-31-13	1	7 entries
3	VH3-11-1	0	
3	VH3-11-2	0	
3	VH3-11-3	5	
3	VH3-11-4	0	
3	VH3-11-5	1	
3	VH3-11-6	1	
<b>3</b> .	VH3-11-7	0	
3	VH3-11-8	5	
3	VH3-13-1	9	
3	VH3-13-2	3	
3	VH3-13-3	0	
3	VH3-13-4	0	•
3	VH3-13-5	0	•
3	VH3-13-6	0	
3	VH3-13-7	32	
3	VH3-13-8	4	
3	VH3-13-9	0	
3	VH3-13-10	46	
3	VH3-13-11	0	
3	VH3-13-12	11	
3	VH3-13-13	17	•
3	VH3-13-14	8	
3	VH3-13-15	4	
. 3	VH3-13-16	3	
3	VH3-13-17	2	
3	VH3-13-18	1	
3	VH3-13-19	13	
3	VH3-13-20	1	
3	VH3-13-21	1	
3	VH3-13-22	0	

Table 3C: (continued)

Family <sup>1</sup>	Name	Rearranged <sup>2</sup>	Sum
3	VH3-13-23	0	
3	VH3-13-24	4	
3	VH3-13-25	1	
3	VH3-13-26	6	
3	VH3-14-1	1	
3	VH3-14-4	15	
3	VH3-14-2	0	•
3	VH3-14-3	0	
3	VH3-1X-1	0	
3	VH3-1X-2	0	•
3	VH3-1X-3	6	
3 3	VH3-1X-4	0	•
3	VH3-1X-5	0	
3	VH3-1X-6	11	
3	VH3-1X-7	0	
3	VH3-1X-8	1	
3	VH3-1X-9	0	212 entries
4	VH4-11-1	0	
4	VH4-11-2	20	
4	VH4-11-3	0	
· <b>4</b>	VH4-11-4	0	
4	VH4-11-5	0	
4	VH4-11-6	0	
4	VH4-11-7	5	
4	VH4-11-8	7	
4	VH4-11 <b>-9</b>	3	
4	VH4-11-10	0	•
4	VH4-11-11	0	
4	VH4-11-12	4	
4	VH4-11-13	0	
4	VH4-11-14	. 0	
4	VH4-11-15	0	•
4 .	VH4-11-16	1	
4	VH4-21-1	0	
4	VH4-21-2	0	
4	VH4-21-3	1	
4	VH4-21-4	1	

Table 3C: (continued)

Comital	Nomo	Poorrong 2	
Family <sup>1</sup>	Name	Rearranged <sup>2</sup>	Sum
4	VH4-21-5	1	
4	VH4-21-6	1	
. 4	VH4-21-7	0	
4	VH4-21-8	0	
. 4	VH4-21-9	0	
4	VH4-31-1	0	
4	VH4-31-2	0	
4	VH4-31-3	0	
4	VH4-31-4	2	•
4	VH4-31-5	0	
4	VH4-31-6	0	
4	VH4-31-7	0	
4	VH4-31-8	0	
<b>4</b> .	VH4-31-9	0	
4	VH4-31-10	0	
4	VH4-31-11	0	
4 ′	VH4-31-12	4	
4	VH4-31-13	7	
4	VH4-31-14	0	•
4	VH4-31-15	0	
4	VH4-31-16	0	
4	VH4-31-17	. 0	. •
4 .	VH4-31-18	0	
4	VH4-31-19	0	
4	VH4-31-20	0	57 entries
5	VH5-12-1	82	
5	VH5-12-2	1	
5	VH5-12-3	0	
5	VH5-12-4	14	97 entries
6	VH6-35-1	74	74 entries

Table 4A: Ana	lysis of V	' kappa	subgroup	1
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4A: Analysis of v	Framework I															
amino acid'	<del>-</del>	2	က	4	2	9	7	8	6	10	11	12	13	14	15	16
Α		1							1				102		1	
В			1			1							<u> </u>		<u></u>	
С														1	<u>.</u>	
D	64														<u></u>	<u></u>
E	8		14										<u>.</u>		1	<u></u>
F									1	6				1		<u></u>
G			*******					***********					P+10000000000			105
Н																
· ,		65			******										4	:
K			1													
L		6	*******	21				•••••			96		1			: : : : : : : : : :
M	1			<b>6</b> 6												
N			*********		10047000											
Р								103		1		2			1	
Q			62			88					1					
R			*******					•••••								
5							<b>8</b> 9		102	80		103		103		
T		1			88					18						
V		1	9								8		2		98	
W														************		
X	1															
Y																
_												**********				
unknown (?)																
not sequenced	31	31	18	18	17	16	16	2	1							
sum of seq <sup>2</sup>	74	74	87	87	88	89	89	103	104	105	105	105	105	105	105	105
oomcaa3	64	65	62	66	88	88	89	103	102	80	96	103	102	103	98	105
mcaa <sup>4</sup>	D	ı	Q	М	T	а	S	Р	S	S	L	S	Α	S	V	G
rel. oomcaas	%98	%88	71%	76%	100%	%66	100%	100%	%86	76%	91%	%86	97%	%86	93%	100%
pos occupied6	4	5	5				1	1	3	4	3		:			

Table 4A: Analysis of V kappa subgroup 1

	-					-										
amino acid	1 . 1	<u> </u>	8	19	20	21	22	23	24	25	26	27	۷	8	ပ	٥
А				1	1		1			103						
В												1				
. C								105								
D	1	01														
E		2							1	1		2		***********		*********
F					***************************************	2			****							
G											1	******	****			
H												1	************			
				6	4	101	1							**********		
K						•••••••••••••••••••••••••••••••••••••••			2		**********	1	•••••	*******		
L									1				*********	***********		
M																
N ·									***************************************		1		*******	*********		
Р											***************************************			******		
Q									20			100		•••••		
R :			94						81					***********		
S			5		1						102					
Т			6		99		103			1	1			*******		
<u> </u>				98		2							••••••			
W										,			************	*******		
X		1											•••••	•••••		
Y		1														
												-	105	105	105	105
unknown (?	2)										*******		***********			
not sequence	ed													-		
sum of seq	² <u>1</u>	05	105	105	105	105	105	105	105	105	105	105	105	105	105	105
oomcaa3	1	01	94	98	99	101	1.03	105	81	103	102	100	105	105	105	105
mcaa <sup>4</sup>		D	R	V	T	ı	T	С	R	Α	S	Q	-	_	-	_
rel. oomcaa	35	<b>96</b> %	%06	93%	94%	%96	%86	100%	77%	%86	97%	95%	100%	100%	100%	100%
pos occupie	q <sub>e</sub>	4	3	3	4	3	3	1	5	3	4	5	1	1	1	1

WO 97/08320 .

Table 4A: Analysis of V kappa subgroup 1

•	CDRI														
amino acid¹	ш	ய	28	29	30	31	32	33	34	35	36	37	38	39	40
Α					1	1		1	42						
В							•					1	1	, , , , , , , ,	
. C							1								
D			25		1	5	7					1			
E							1			***********		2			
F				1	1		7	**********	******		6				
G			25		7	3	•••••		4						
H		,			1	2	2	••••	1			2			
1		••••••••		98	1	4			1		************				
K						7		*********				•••••		95	
L					2	1	*********	101	***********		••••				
M										-	·~	*******			
N			6		16	42	*********	*******	50						
. Р							**********	******	••••••	****					102
Q											*********	98	103	2	
R					16	3	2	•••••	******	•••••	•••••			3	1
S			41	2	57	32	3	1	1	••••					1
T			7			4	•••••	•••••	4					1	
V			1	4	1		••••••	1	***********	***********	F P			•••••	
W							21	*****	••••••	104			·····		
Χ			······						1	••••••					
Υ					1		60				98				
	105	105					•••••	••••••	*********	********	******				
unknown (?)				.,		***********								3	
not sequenced						1	1	1	1	1	1	1	1	1	1
sum of seq²	105	105	105	105	105	104	104	104	104	104	104	104	104	104	104
oomcaa <sup>3</sup>	105	105	41	98	57	42	60	101	50	104	98	98	103	95	102
mcaa <sup>4</sup>	-		S		S	N	Y	L	N	W	Υ	Q	Q	Κ	Р
rel. oomcaa³	100%	100%	39%	93%	54%	40%	58%	97%	48%	100%	94%	94%	%66	91%	%86
pos occupied <sup>6</sup>	1	1	6	4	12	11	9	4	8	1	2	5	2	4	

Table 4A: Analysis of V kappa subgroup 1

	Fran	newor	k II										DR II		
amino acid'	41	42	43	44	45	46	47	48	49	20	51	25	53	54	55
Α			94							50	95				
В					-										
. C															
D										21	1	1	1		*********
E	1	3			1	1				. 1		1			33
F						1			3			. 1			
G	100		1							9	2				
Н									2						1
		1				1		100					1		
K		95			86					16			2		5
L		1				89	103							101	
M								2							
N					10					2		1	25		
Р				104						1					1
Q		1			1										62
R					3	3	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,			·············			1	1	2
S					1				5	1	1	99	41	2	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,
Ţ		3			1					1	4	1	31		
V			9			9					1		1		******
W													************		
X					1						••••		1		
Y									92	1					
-											•••••		********		
unknown (?)	3				••••	**********					*****		*********		
not sequenced	1	1	1	1	1	1	2	3	3	2	1	1	1	1	1
sum of seq <sup>2</sup>	104	104	104	104	104	104	103	102	102	103	104	104	104	104	104
oomcaa³	100	95	94	104	86	89	103	100	92	50	95	99	41	101	62
mcaa <sup>4</sup>	G	Κ	Α	Р	K	L	L	1	Υ	Α	Α	S	S	L	Q
rel. oomcaa <sup>s</sup>	%96	91%	%06	100%	83%	96%	100%	%86	%06	49%	91%	95%	39%	97%	60%
pos occupied	}		:	<u> </u>	:	:	•	•	:	:		:	:	:	:

		r –													
amino acid'	56	57	58	59	09	61	62	63	64	65	99	29	89	69	70
Α	3										2	1	1	1	
В				1											
. C								,							
D	1														67
E	·						,						1		30
F			1				103				•	3			
G	2	105							105	4	101		102		
Н	<u></u>	)	• • • •	• • • •			<b>)</b>								3
	3	***********	4	• • • • •			1	3							
K	1				•••••••	1		*********	***********		•••••	: : : : :		: : : :	1
L		••••••••						1	••••••	***********	***************************************	: : : : : : ;		• • • • • • • • • • • • • • • • • • •	
M								•••••		*********	**********			1	
N	6	•••••				••••••									
P	1			101	2					•••••	************	************************	••••••••		
<u> </u>		********		•••••				••••••		1	•••••				
R	1			********	**********	103		1		1	1		***********	2	
S	68			· 2	103	•••••		98		96		100			
T	19		•	1		1		2		3		**********	*****	101	·
V			<b>9</b> 9			*********	1		************			•••••	•••••		1
W			•••••			•					**********			•••••	
Χ			1		••••••						1	************	1	********	2
Y												1			1
			*********	•••••••	**********	•••••••••••••••••••••••••••••••••••••••					***************************************	***********		*********	
unknown (?)			*****		*************		************						***************************************	•••••	
not sequenced															
sum of seq²	105	105		:	7			•	105	105	105	105	105	105	105
oomcaa <sup>3</sup>	68	105	99	101	103	103	103	98	105	96	101	100	102	101	67
mcaa'	S	G	V	Р	S	R	F	S	G	<u>S</u>	G	S	G	T	D
rel. oomcaa <sup>s</sup>	65%	100%	94%	<b>%96</b>	98%	98%	986	93%	100%	91%	%96	95%	97%	<b>%96</b>	64%
pos occupied <sup>6</sup>	10	1	4	4	2	3	3	5	1	5		:	:		:::::::::

Table 4A: Analysis of V kappa subgroup 1

A. Allalysis Of		amev													
amino acid¹	71	72	73	74	75	92	77	78	62	80	81	82	83	84	82
Α		3		İ		1				2				101	1
В					1				3		2				
. C															
D						1					16	101			
E											83				*************
F	102	1	21										73		
G							4				1			2	
Н				······											
1					99	. 5						***************************************	17		
K												.,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,			
L			81					103	1				1		
М															1
N			************************			7	4				***************************************				1
Р										97					1
Q									97						
R						2	1		2		••••••				
S		2		1		86	94			4			1		
T		98		102		2	1						•••••		97
V	1		2		4			1					11	••••••	1
W											***************************************				
X				1			.,				1	2	••••		
Y	1														
-					400 100000		•••••				•••••	•••••••			
unknown (?)													••••••		
not sequenced	1	1	1	1	1	1	1	1	2	2	2	2	· 2	2	3
sum of seq²	104	104	104	104	104	104	104	104	103	103	103	103	103	103	102
oomcaa <sup>3</sup>	102	98	81	102	99	86	94	103	97	97	83	101	73	101	97
mcaa*	F	T	L	Τ	1	S	S	L	Q	Р	E	D	F	Α	T
rel. oomcaa <sup>s</sup>	%86	94%	78%	%86	95%	83%	%06	%66	94%	94%	81%	%86	71%	986%	95%
pos occupied <sup>s</sup>	3	4	3	:	:	7	5	:	:	3	5	2	5	2	

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Table 4A: Analysis of V kappa subgroup 1

4A. Alialysis of										CDR I	II	<del></del>				
amino acid'	98	87	88	68	06	91	92	93	94	95	4	æ	U	۵	ш	<u>.</u>
Α					1	7	1		5	1						
В				2	3					·					<u> </u>	
. C			102									1.				
D							23	5	1							
E							1	1		1	1					
F		7				3			13							
G						1		1	2	1		1				
Н		1		4	6	7	3	1								
							4	1	2	1						
K	1				7		1									
L				7		6	2		18	2						
M													,			
N					**********	6	31	19	1							
Р									1	82	6					
Q		***********		90	86	1	2									
R		************				1		2	2	************						
S	1	************				27	3	58	5	10						
T		-				3	1	15	25							·
V		***************************************							5							
W					**********				1							
X		**********												,		
Υ	101	93				42	32	1	23							
-		********						*************		3	82	88	89	89	89	89
unknown (?)		1														
not sequenced	2	3	3	2	2	1	1	1	1	4	16	16	16	16	16	16
sum of seq²	103	102	102	103	103	104	104	104	104	101	89	89	89	89	89	89
oomcaa³	101	93	102	90	86	42	32	58	25	82	82	88	89	89	89	89
mcaa•	Υ	Υ	С	Q	۵	Υ	Υ	S	T	Р		-	-	-	-	-
rel. oomcaa <sup>s</sup>	%86	91%	100%	87%	83%	40%	31%	26%	24%	81%	92%	%66	100%	100%	100%	100%
pos occupied <sup>6</sup>	3	3	1	4	•	:	12	•	14				•	1	1	1

Table 4A: Analysis of V kappa subgroup 1

.,		_	<u>'</u>				Fra	mev	vork	iV					
amino acid'	96	97	86	66	100	101	102	103	104	105	106	⋖	107	108	sum
Α	1														627
В					1					1					19
. C															209
D	1									15					459
E					2					65					258
·F	6		86							***********	2				451
G				87	29	87								2	894
Н	2	1													40
<b></b>	5								1		72				606
Κ	1	1						77			********		79		480
<u> </u>	18	1	1						22	4	2				793
M		1									5				77
N	1										1		2		232
Р	6				7									1	620
Q	1			******	48					1					865
R	6		••••••	*****				6					2	70	413
S	2	2			*******			•••••							1636
T	2	82					87	3					2		1021
V	2			*******				1	63		3				440
W	15			•••••											141
X		•••••	••••••	•••••				*******	••••••	•••••				•••••	14
Υ	16														564
-	4	1		•••••			•••••	*******		•••••		85	******	1	1250
unknown (?)				•••••				••••••		••••••		********	••••		7
not sequenced	16	16	18	18	18	18	18	18	19	19	20	20	20	31	589
sum of seq <sup>2</sup>	89	89	87	87	87	87	87	87	86	86	85	85	85	74	
oomcaa <sub>3</sub>	18	82	86	87	48	87	87	77	63	65	72	85	79	70	
mcaa <sup>4</sup>	L	T	F	G	G	G	Ţ	K	٧	E	l	-	K	R	
rel. oomcaas	20%	92%	%66	100%	55%	100%	100%	9,68	73%	9097	85%	100%	93%	95%	**************************************
pos occupied6	17	7	2	1	5	1	1	4	3	5	6	1	4	4	

Table 4B: Analysis of V kappa subgroup 2

		•									Frar	new	ork	ı							
amino acidi	-	7	က	4	5	9	7	8	6	10	11	12	13	14	15	16	17	18	19	20	21
А																:			22		
В																					
· C																					
D	14																				
E	3					9											15				
F									1	1											
G									••••••							22					
Н									•••				•••••	· •		: : :					
ļ ·		8					********												••••		22
K		•••••																			
L		3		1					17		18				6						
M		•••••		15																	
N																					
Р		••••						18				18			15			22			
Q						18						<u></u>					7				
R																•••••					
S							18			17										22	
T					17									21							
V		6	17	1									18								
W																					
X																					
Υ																					
unknown (?)					1																
not sequenced	5	5	5	5	4	4	4	4	4	4	4	4	4	1	1						
sum of seq²	17	17	17	17	18	18	18	18	18	18	18	18	18	21	21	22	22	22	22	22	22
oomcaa <sub>3</sub>	14	8	17	15	17	18	18	18	17	17	18	18	18	21	15	22	15	22	22	22	22
mcaa*	D	•••••••••••••••••••••••••••••••••••••••	٧	•••••••	Ţ	•••••••••••••••••••••••••••••••••••••••	····÷	•••••••••••••••••••••••••••••••••••••••	····÷		••••••	Р	•••••••••••••••••••••••••••••••••••••••	Т	···	G	•••••••••••••	Р	Α	S	1
rel. oomcaa <sup>s</sup>	82%	47%	100%	%88	94%	100%	100%	100%	94%	94%	100%	100%	100%	100%	71%	100%	%89	100%	100%	100%	100%
pos occupied <sup>6</sup>	2	3	1	3	1	1	1	1	2	2	1	1	1	1	2	1	2	1	1	1	1

·					<u>:</u>				•		CDF	RI	· · · · · ·	-						Ī	
amino acid'	22	23	24	25	26	27	∢	8	U	۵	ш	u	28	29	30	31	32	33	34	35	36
Α																					
В																<b></b>		<b></b>	<b></b>		•••••
· C		22																			
D	<b>.</b>	<u></u>	<u></u>							1			9		1	1			11		
E		<u>.</u>	<u>.</u>																		
F				<u></u>											2						7
G	<b>.</b>	<u>:</u> : : :									1			22							
Н	<b>.</b>									16							1		1		
ı																				•••••	
K			1													1					
L						1		22	13									22			
M									1												
N													10		7	12			9		
Р																					
Q	1		******			21															
R			21								2	·						·			
S	21			22	22		22				19		1								
T			•••••		•••••											8					
V					******				8												
W			•••••		••••••		******			1					,					22	
X											,		1		1				1		
Υ										4			1		11		21				15
-												22									
unknown (?)						•••••										••••					
not sequenced																					
sum of seq <sup>2</sup>	22	22	22	22	22	22	22	22	22	22	22	22	22	22	22	22	22	22	22	22	22
oomcaa¹	21	22	21	22	22	21	22	22	13	16	19	22	10	22	11	12	21	22	11	22	15
mcaa'	S	С	R	S	S	Q	S	L	L	Н	S	-	N	G	Υ	N	Υ	L	D	W	Υ
rel. oomcaa <sup>s</sup>	95%	100%	95%	100%	100%	95%	100%	100%	29%	73%	%98	100%	45%	100%	20%	55%	95%	100%	20%	100%	%89
pos occupied <sup>6</sup>										:			:		:		•••••••••••••••••••••••••••••••••••••••	1	•	•	2

Table 4B: Analysis of V kappa subgroup 2

ie 40. Alialysis ul		_	-	<u>-</u>	Fran		ork	11								(	CDR	11		•	
amino acid'	37	38	39	40	41	42	43	44	45	46	47	48	49	20	51	52	53	54	55	26	57
Α																			14		
В		•																			
· c		<u> </u>			<u></u>	<u></u>	<u></u>														
D	ļ	<u>.</u>	ļ		<u></u>		<u></u>	<u></u>		<u></u>		: :							7		
E	<b></b>	<u></u>		<u></u>	<u></u>		<u></u>	<u></u>	1	<u></u>	<u></u>	<u></u>	<u></u>	<u>.</u>							
F	<b></b>	<u></u>	<u></u>	<u>.</u>	<u> </u>			<u>.</u>	<u></u>	<u> </u>	<u></u>		<u>.</u>		,						
G	<b></b>				22			<u></u>		<u></u>	<u></u>		<u></u>		12				1		22
Н					<u></u>			<u>.</u>													
1	ļ									1		22									
K		<u></u>	15									<u></u>		5							
L	16							·		14	21			14	1	•••••					
M								<u> </u>													
N						••••			***************************************								18				
Р		••••••		22		•••••		21	.,,,,,,,,		••••••										
Q	6	22	••••••			22			12	•••••	•••••	••••		1							
R			7						8	7				1				22			
<u>S</u>		••••••			·····		21		••••••	••••••		••••••	•••••		2	22	2			22	
T		•••••		*********					••••••			•••••					1				
<u> </u>			••••••							********	1				6						
W				••••••			••••••		••••••		*********										
X				•••••	•••••																
Υ											-		21				1				
-				•••••					******												
unknown (?)		•••••	********	-1000444		•••••	• •••••••		••••••	••••••			••••••								
not sequenced									1	_			_	1	_						_
												22		:	•••••••	•••••••••••••••••••••••••••••••••••••••	•••••••••••••••••••••••••••••••••••••••	•••••••••••••••••••••••••••••••••••••••	•••••••••••••••••••••••••••••••••••••••	•••••••	
oomcaa,									12	14	21	22	21	14	12	22	18	22	14	22	22
mcaa¹	•••••••••••••••••••••••••••••••••••••••	Q	••••••	Р	•••••••••••••••••••••••••••••••••••••••	Q			Q	L	L	ı	Υ	••••••	G	•••••••••••••••••••••••••••••••••••••••		R	•••••••••••••••••••••••••••••••••••••••	S	G
rel. oomcaa <sup>s</sup>	73%	100%	%89	100%	100%	100%	100%	100%	27%	64%	95%	100%	100%	%29	57%	100%	82%	100%	64%	100%	100%
pos occupied <sup>a</sup>	2	1	2	1	1	1	1					1	:	:	:	•	•		3	:	·····

														Fra	me	wor	k III				
amino acid'	28	59	09	61	62	63	64	65	99	29	89	69	70	7.1	72	73	74	75	9/	77	78
Α																					
В										: :				}							<b>••••••••</b>
· C														************						•••••	
D			22				1				1		22							•••••	
E														*******	••••••	•	•••••			•••••	
F					21	••••••	••••••	••••••••••••••••••••••••••••••••••••••	•••••	•••••	••••••		•••••	22	••••	•••••	•••••	••••			
G						••••••	21		22		21	•••••				••••••					
Н		• • • • • • • • • • • • • • • • • • •				••••••			•••••		••••	•••••		•••••			•••••	•••••			
1										•	•••••		•••••	•••••	,		1	21			
K												******			*******	•••••	19		•••••		
L											•••••	••••		_		21	1				
M										•		********	••••••			•	••-		••••	••••••	******
N									·											······•	*****
Р		22							**********		***************************************	*******		******			•				******
Q														********	••••••					•	******
R			·	20				1				••••••			-	***********				20	***************************************
S				1		22		21		22	*******					***************************************	******		20	1	
T				1								22	•••••••		21	********			1	•••••••••••••••••••••••••••••••••••••••	
V	22				1							•		•••••••							21
W																***************************************			·		*******
X														•							******
- Y																********	•••••				*******
unknown (?)					,			•••••							1						
not sequenced																1	1	1	1	1	1
sum of seq²	22	22	22	22	22	22	22	22	22	22	22	22	22	22	22	21	21	21	21	21	21
oomcaa <sup>3</sup>	22	22	22	20	21	22	21	21	22	22	21	22	22	22	21	21	19	21	20	20	21
mcaa•	٧	Ρ	D	R	F	S	G	S	G	S	G	Τ	D	F	T	L	Κ	١	S	R	٧
rel. oomcaas	100%	100%	100%	91%	95%	100%	95%	95%	100%	100%	95%	100%	100%	100%	95%	100%	%06	100%	95%	95%	100%
pos occupied <sup>a</sup>		1		3							:	:	1	:				1		•	1

Table 4B: Analysis of V kappa subgroup 2

e 45: Analysis 0				- 3													(	DR	111		
amino acid'	79	80	.81	82	83	84	85	98	87	88	88	90	91	92	93	94	95	۷	8	ပ	۵
A		20											14			1					
В		<u>.</u>	<u></u>	<u></u>								1			1						
· C	<b></b>									21											
D	<b>.</b>		1	21	<u>.</u>					<u></u>											
E	19		20					: : :	<u></u>												
F	<b>.</b>				<u></u>		<u>.</u>	<u>.</u>	<u></u>	<u></u>											
G	1					21							6			1		2			<b></b>
Н													1		7						
	ļ						1							••••		1					
K																					
<u> </u>		•••••	•••••				1							12			2				
M											21										
N													••••								
Р		1	•••••	r*******	••••••			•••••		•••••						2	16	1			
Q	1											20			13						
R						•••••		•••••						1							
<u>S</u>		•••••	•••••		•••••	••••••		•••••			•••••	,				3	2				
T		.,,,,,,,			••••	•••••				••••••	•••••			8		7					
V					21		19														
<u> </u>					•••••	•	•••••									6		••••••			
X				•				•••••		••••••	•••••										
Y								21	21						_						
						•••••			••••••									14	17	17	17
unknown (?)						•••••				••••••											
not sequenced		1				_			1	_		_			1	=					_
sum of seq <sup>2</sup>												:	•••••••••••••••••••••••••••••••••••••••	•••••••••••••••••••••••••••••••••••••••	•••••••••••••••••••••••••••••••••••••••					••••••	•••••••
									*********			20		•••••••••••••	••••••	•••••••••••••••••••••••••••••••••••••••	•••••••••••••••••••••••••••••••••••••••		17	17	17
mcaa <sup>4</sup>	Ε	Α					*******				••••••	Q	•••••••••••••••••••••••••••••••••••••••	•••••••••••••••••••••••••••••••••••••••	•••••••	•••••••••••••••••••••••••••••••••••••••			-	-	-
rel. oomcaa <sup>s</sup>	<b>%06</b>	95%	95%	100%	100%	100%	<b>%</b> 06	100%	100%	100%	100%	95%	%29	57%	62%	33%	%08	82%	100%	100%	100%
pos occupied <sup>6</sup>	3	2	2	1	1	1	3	1	1	1	1	2	3	3	3	7	•			•	•

Table 4B: Analysis of V kappa subgroup 2

									Fra	me	wor	k IV					
amino acid'	ш	ц.	96	97	98	66	100	101	102	103	104	105	106	٧	107	108	sum
Α																	71
В				••••••			••••••	•	•••••	•••••		1		•••••	•••••	·····	3
С												•••••			*******	•••••	43
D																	112
<u>E</u>				*******			•••••					13					71
. F			1		17	*****	••••										72
G						17	2	16				1		·			233
Н			•••••	•••••			•••••										26
			3										14				94
K		,								12					13	·	66
<u>L</u>			2								11						219
М																	37
N																	56
Р			1														159
Q			1				14									*****	159
R										4						12	126
<u>S</u>																	325
T				17					16								140
V											5						146
W			2														31
X																	3
Y			7														123
_	17	17												13			134
unknown (?)				•••••													2
not sequenced	5	5	5	5	5	5	6	6	6	6	6	7	8	9	9	10	211
sum of seq <sup>2</sup>	17	17	17	17	17	17	16	16	16	16	16	15	14	13	13	12	
oomcaa,	17	17	7	17	17	17	14	16	16	12	11	13	14	13	13	12	
mcaa <sup>4</sup>	-	-	Υ	T	F	G	a	G	T	K	L	Ε	ı	-	Κ	R	
rel. oomcaas	100%	100%	41%	100%	100%	100%	988%	100%	100%	75%	%69	87%	100%	100%	100%	100%	
pos occupied <sup>6</sup>	1	1	7	1	1	1	2	1	1	2	2	3	1	1	1	1	

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Table 4C: Analysis of V kappa subgroup 3

	· Framework I															
amino acid'	-	7	r.	4	5	9	7	æ	6	10	11	12	13	14	15	16
Α		5					2		27						1	
В	1															
· C												2				
D	2								14							
E	76	*********	27													
F .		1		<u> </u>										1		
G	1								82						1	152
Н		**********								1						
		75														
K	3															
L		4	1	104			1				150		129		1	
·M	5			13												
N														5		
Р								124				•			147	
Q				·		123										
R					1											
S							119		3	1		150	1	141		
<b>T</b>		2			117			-		147				5	1	•
V		1	89	1			1				1		22		. 1	
W																
X	i				-											
Υ																
unknown (?)								·		,						
not sequenced																
sum of seq'	88	88	117	118	118	123	123	124	126	149	151	152	152	152	152	152
oomcaa,	76	75	89	104	117	123	119	124	82	147	150	150	129	141	147	152
mcaa*	Ε	1	٧	L	T	Q	S	Р	G	T	L	S	L	S	Р	G
rel. oomcaas	%98	85%	76%	9/88	%66	100%	97%	100%	65%	%66	%66	%66	85%	93%	97%	100%
pos occupied <sup>6</sup>	6	6	•••••••••••••••••••••••••••••••••••••••					1					:	:		

Table 4C: Analysis of V kappa subgroup 3

C +C. / mary 5/5 0/		' <del>-</del>				· · · ·	-	T								CDRI
amino acid'	17	18	19	20	21	22	23	24	25	26	27	⋖	æ	U	۵	ш
Α			178	2		:			166	1						
В								•				•	•			
. С			•		<del></del>		181	†*************************************	•	1			<u> </u>			
D	6						•		<u>.</u>			<del></del>	<u> </u>		<u> </u>	
E	146	1				•		<del></del>	<b></b>		1	•••••••••••••••••••••••••••••••••••••••	<u> </u>		<u> </u>	
F ·					7	1	**************************************			•		•••••••••	•			
G	1	1							7	1	÷	1				
Н		• • • • •									17		•	<u></u>		<b></b>
1		1		5	2				• • • • • • • • • • • • • • • • • • •		•				•	
K		1						5								
L					173						1	1		<u>.</u>		
·M												**************************************	 :			
N												9				
Р																
Q											159					
R		175	************		•••••			176		1	1	10				
5						180	• • •	-	7	175		87				
T		1		174	******				7	2		1				
V		1	4	1					1			1				
W			***********					1								
X																
Y						1					1					
					***************************************					***********		72	182	182	182	182
unknown (?)					***********			•••••		•••••	1	•	***********	••••		
not sequenced																
sum of seq <sup>2</sup>	153	181	182	182	182	182	181	182	182	181	181	182	182	182	182	182
oomcaa <sup>,</sup>	146	175	178	174	173	180	181	176	166	175	159	87	182	182	182	182
mcaa•	Ε	R	Α	T	L	S	С	R	Α	S	Q	S	-	-	-	-
rel. oomcaa <sup>s</sup>	95%	97%	%86	%96	95%	%66	100%	92%	91%	97%	88%	48%	100%	100%	100%	100%
pos occupied <sup>6</sup>	3	7				3	1	3	5					1	1	1

									<u> </u>						Fra	mew
amino acid'	и.	28	29	30	.31	32	33	34	35	36	37	38	39	40	4	45.
Α				1	1			181								
В																
. · C				•												
D			1	1	2	1									!	
Е						1							1			
F		1				7				1				***************************************	•	••••••••••••••••••••••••••••••••••••••
G			2	7	3	1		2			<u>.</u>	<del></del>	<u></u>	1	184	
Н			1			2				1	<u> </u>	12	1	1		
		24	4	1	1									•		
K				1	1								153			
L		8	1			1	176					3		• • • • • • • • • • • • • • • • • • •		
·M		-														
N			3	12	25	32										
Р					1									170		•••••
Q					1	1					183	167	1			18
R			10	3	18	16		1			1		27	5	•	•
· S		72	86	151	118	4								5		
T		1	1	3	8	1							1	,		
<u>V</u>		76	68		1	-	7					3		2		
W			5						185							
X						**********										
Y				1	1	115				183						
	182		,													*****
unknown (?)						•	**********				1					•••••
not sequenced																
sum of seq <sup>2</sup>	182	182	182	181	181	182	183	184	185	185	185	185	184	184	184	184
oomcaa,	182	76	86	151	118	115	176	181	185	183	183	167	153	170	184	18
mcaa'	_	٧	S	S	S	Υ	L	Α	W	Υ	Q	Q	K	Р	G	Q
rel. oomcaa <sup>s</sup>	100%	42%	47%	83%	65%	63%	%96	%86	100%	%66	%66	<b>20%</b>	83%	92%.	%00I	980%
pos occupied <sup>6</sup>	1			10	•••••••••••••••••••••••••••••••••••••••			:		3	:	:	:	:		

+C. / ilidiy5i5 01	rk II									-	CDR	11				
amino acid¹	43	44	45	46	47	48	49	20	51	52	53	54	55	26	57	28
Α	176							4	147				176	1		
В										<u></u>	<b>.</b>		<del> </del>			<del></del>
· C									1			**************************************				
D							·	43					2		4	
E															·	
F.				1		1	4							•		
G								125					2	10	179	
Н							9		1							
<u> </u>						178								1		168
K			1								7	1				
L		1		179	174	1										
· M						3					1					
N.			1					1			53		·	2		
Р	5	184								2			2	2		
Q							1									
R			182					1			4	180				
S		·		٠.			3	6	4	179	74	1		5		-
T	3								11	2	44			164		2
V				3	. 9		-	3	19				3			15
W							1					1				
X																
Υ							165		·						2	
••																
unknown (?)			1													
not sequenced																
sum of seq <sup>7</sup>	184	185	185	183	183	183	183	183	183	183	183	183	185	185	185	185
oomcaa,	176	184	182	179	174	178	165	125	147	179	74	180	176	164	179	168
mcaa*	Α	Р	R	L	L	ı	Υ	G	Α	S	S	R	Α	Τ.	G	١
rel. oomcaa'	%96	%66	98%	98%	95%	9/0/6	%06	9/89	80%	%86	40%	98%	95%	%68	97%	91%
pos occupied"	3	2	3											***********		3

													F	rame	work	: 111
amino acid	59	09	61	62	63	64	65	99	67	89	69	70	71	72	73	74
A		68	3					3	3		3	3 1		3		
В													•			•
. C									<u> </u>			-				·
D		112				1	•		<u> </u>	<u> </u>		152		÷	÷	<del></del>
Е	·		•	•			•	1		1	<u> </u>	30	<u> </u>	<b></b>	<del>•</del>	<u> </u>
F.				183	<b>†</b>		***********				·	<u> </u>	183	<b></b>	2	
G						184	3	178	_	177	<u> </u>	÷	<del></del>	<del> </del>	<u> </u>	<del></del>
Н		1								:		÷·····	<u> </u>		<b></b>	
1				1					:		**********	**********	•	1	••••••••••••••••••••••••••••••••••••••	3
K	9 8 8	<u> </u>	1								<u> </u>				<u>.</u>	
L				1						<del></del>	÷	••••••••••••••••••••••••••••••••••••••	•		182	
. M								1	•		••••••••••••••••••••••••••••••••••••••	¢				
N		1											***********	1		
Р	177								÷		············	•			***********	
Q									• · · · · · · · · · · · · · · · · · · ·	• ····································	••••••••	1				
R			182		2		1				2		*********		••••••	
S	` .7				180		179	·	185		3			7		2
T	1		2		3	**********	2	·			177	·		172		179
V		3					•	1		1						
W					*********		-			1						
X		*********												•••••••••••••••••••••••••••••••••••••••		
Y													1			
							**********									
unknown (?)			*********					1								
not sequenced																
sum of seq?	185	185	185	185	185	185	185	185	185	185	185	184	184	184	184	184
oomcaa³	177	112	182	183	180	184	179	178	185	177	177	152	183	172	182	179
mcaa'	Р	D	R	F	S	G	S	G	S	G	T	D	F	T	L	T
rel. oomcaas	<b>%96</b>	61%	%86	%66	92%	%66	92%	%96	100%	<b>%96</b>	%96	83%	%66	93%	%66	97%
pos occupied <sup>a</sup>	3	5	············	:		······································		5	:	:	•	:	2	5 5	2	
							1 1	0	************							;

amino acid'	75	9/	11	78	79	80	81	82	83	84	85	98	87	88	83	90
А							3			174						
В					1		**************************************			• • • • • • • • • • • • • • • • • • •	••••••••••••••••••••••••••••••••••••••	<b></b>			••••••••••••••••••••••••••••••••••••••	
· C		• • • • • • • • • • • • • • • • • • •					•		2			<u> </u>	1	182	·	
D			1				3	182								
E					149		175				•	• • • • • • • • • • • • • • • • • • •			# ************************************	2
F		1							178		2	1	4	• • • • • • • • • • • • • • • • • • •		
G			3					1		2						
Н				•••••	***********						1				1	7
	178				*********			1	1		9					
K							1									
L				178		1			1		7		1			1
M						•				1	5					
N	1	5	***********													
Р						149	•				•		•••••			
Q					34									1	181	155
R		1	111							3	)					1
S		169	65			34			1				2			
T		. 8	4							1						8
V	4			6					1	3	159					7
<u> </u>												•••••				
X		••••					•••••									
Υ	1										1	183	176		1	2
-		•••••														····
unknown (?)				•••••	•••••								••••••			
not sequenced																
sum of seq <sup>2</sup>							*************		•••••••		***********		**********	***************************************	••••••	
oomcaa <sup>3</sup>	178		111	178			175	***************************************	178			183	176	182	181	155
mcaa⁴	1	S	R	L	Ε	Р	E	D	F	Α	V	Υ	Υ	С	Q	Q
rel. oomcaa <sup>s</sup>	92%	92%	<b>%</b> 09	97%	81%	81%	%96	%66	92%	95%	96%	%66	%96	%66	%66	85%
pos occupied <sup>6</sup>	4	5	5	2	3	3		3			7	2	5	2	3	8

Table 4C: Analysis of V kappa subgroup 3

e 4C: Analysis of				· ·		CDR	111									
amino acid'	91	92	93	94	95	⋖	8	ပ	٥	ш	щ.	96	97	86	66	100
Α		1	8	3	3						•					1
В											•					
· c	2			1							•	2	2	-		
D		8	5					<u> </u>				<u> </u>	1		•	
E		2							***************************************	•		1		•		
F .	5	<u>.</u>	2									7		166	•	
G	1	104	15		1	1	2					1			166	41
Н	4	1										2		: :		:
1		· • • • • •	1			1						4				
K	ļ		2		<u>.</u>	1			•			1		: :		1
L		·········	•	2	7	5						42				
· M		1			1	2										
N		28	71		· · · · · · · · · · · · · · · · · · ·		<u>.</u>	<u>.</u>	<u>.</u>	<u> </u>		1				
Р				1	139	24	<u> </u>	<u>.</u>	<u></u>			7	2			9
Q	1	••••••	1		3	1	<u>.</u>	<u> </u>		<u>.</u>	· · · · · · · · · · · · · · · · · · ·	3				114
R	34	2	3	•••••	2	2	<u>.</u>	<u>.</u>	<u>.</u>	<u>.</u>	<u>.</u>	19				
S	2	33	58	102	15	2		<u> </u>	<b>.</b>	<u>.</u>		1	8			
T		2	13	1	1	·2		· •				1	154			
V					3	• 1	· ·	· ·	<u>.</u>			2				
· W				69				<b></b>	<u>.</u>			24				
X			***************************************			••••••	************		• • • • • • • • • • • • • • • • • • •	**********						
ΥΥ	134	1	1									43				
-			3	3	7	127	167	169	169	169	169	8	1	1	1	1
unknown (?)										•••••						
not sequenced						14	14	14	14	14	14	14	17	16	16	16
sum of seq <sup>2</sup>	183	183	183	182	182	169	169	169	169	169	169	169	166	167	167	167
oomcaa¹	134	104	71	102	139	127	167	169	169	169	169	43	154	166	166	114
mcaa'	Υ	G	N	S	Р	-	-	-	-	<u>-</u> ·	-	Υ	Ţ	F	G	Q
rel. oomcaa <sup>s</sup>	73%	57%	39%	26%	76%	75%	%66	100%	100%	100%	100%	25%	93%	%66	%66	%89
pos occupied <sup>a</sup>	8	11	13	8	11	12			1	1	1	18	:	:	:	:

			F	rame	work	IV					1
	amino acid'	101	102	103	104	105	106	٧	107	108	sum
	Α								Ī		1345
	В										2
	С										375
	D					23					564
	E		•	3		141	<b></b>	<u> </u>	· • · · · · · · · · · · · · · · · · · ·	<del> </del>	759
	F		:		•		6		•	**********	765
	G	166		•	<u> </u>		<u>.</u>	<u> </u>	•	1	1804
	Н					1					64
	i						143	**************************************	<del></del>		803
	K			152					157		489
	L				54		1			2	1596
	М						3				36
	N		1			•			3		255
	Р		1		1						1147
	Q			1		1	*******				1314
ĺ	R			9			2		4	134	1326
	S		2						••••••		2629
	T		162	1	•				1		1593
	V				111		11				646
	W										287
	X										
	Υ			1							1014
	-	1	1	1	1	1	1	166	1	1	2151
	unknown (?)									1	4
	not sequenced	16	16	15	16	16	16	17	17	45	337
	sum of seq'	167	167	168	167	167	167	166	166	138	'
	oomcaa,	166	162	152	111	141	143	166	157	134	
	mcaa⁴	G	T	Κ	٧	E	1	-	Κ	R	
	rel. oomcaa'	%66	97%	%06	0/099	84%	%98	100%	95%	97%	
	pos occupied <sup>a</sup>	2	5	7	4	:	:	1		:	

Table 4D: Analysis of V kappa subgroup 4

							· · · · · · · · · · · · · · · · · · ·				Fra	mew	ork	1				
amino acid'	-	2	က	4	5	9	7	<b>∞</b>	6	10	=	12	13	4	15	16	17	18
Α												24					1	
В																		-
· с										1				***************************************		1		
D	25								26									
E																	25	
F					<u></u>		<u> </u>											
G	<b></b>				<u></u>		<u>.</u>					1				24		-
Н				·			: : :											
1		26	•••••															
K		*****		<u></u>		1												
L				1							26				26			
M			•••••	24		••••												
N	1							-										
Р								26				1						
Q			1			25	••••••	,										
· R																		26
S			•••••				26	**********		25				26		1	·	
T				••••••	26		*********											
V			25	-1									26					
W			*****					*****										
X			*********															
· Y																		
_							********											
unknown (?)																		
not sequenced	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7
sum of seq <sup>2</sup>	26	26	26	26	26	26	26	26	26	26	26	26	26	26	26	26	26	26
oomcaa <sup>3</sup>	25	26	25	24	26	25	26	26	26	25	26	24	26	26	26	24	25	26
mcaa <sup>4</sup>	D		٧	М	Ţ	Q	S	Р	D	S	L	Α	٧	S	L	G	Ε	R
rel. oomcaa <sup>s</sup>	%96	100%	%96	92%	100%	%96	100%	100%	100%	%96	100%	92%	100%	100%	100%	92%	%96	100%
pos occupied <sup>6</sup>	2	1		:		:			••••••			3	•••••••	********	••••••••		2	*******

PCT/EP96/03647

Table 4D: Analysis of V kappa subgroup 4

														CDR	i			
amino acid'	19	20	21	22	23	24	25	26	27	V	8	ပ	٥	m	u.	28	29	30
Α	26						1				1							
В																		
· C					33													
D											1	********	1			1		
E									•					•				
F ·						••••	•	*******	*********			•••••	••••••	•	**********			
G									•			***********	********	••••				
Н						*********	•••••	*******			*********		•••••	•	••••		**********	
1			26	•••••			••••		*********		1	********	*********	•••••	*********		*********	
K					••••••	33	•••••				***********	*******	*******	••••••	•••••	2		30
L ·		•••••									2	_31	********	••••••	•••••			m
· M		•••••						•••••					********	••••••				********
N			************	26	•••••								•••••			30	31	1
Р							1								1			********
Q									32									1
R									1					***************************************			1	1
S .					•		31	33		33		**********	********	32	32		1	*******
T		26											*****	1			••••••	
. <b>V</b>											28	2	**********					
W													*****					
X													***************************************	••••••				*********
Y						-					••••		32		********		•	***********
<b>a</b> as																		
unknown (?)						**********		*********	*********				******	••••••	••••••		*******	*******
not sequenced	7	7	7	7							•••••	•••••	******		*******	•	******	
sum of seq <sup>2</sup>	26	26	26	26	33	33	33	33	33	33	33	33	33	33	33	33	33	33
oomcaa3	26	26	26	26	33	33	31	33	32	33	28	31	32	32	32	30	31	30
mcaa⁴	Α	T	1	N	С	Κ	S	S	Q	5	٧	L	Υ	S	S	N	N	K
rel. oomcaas	100%	100%	100%	100%	100%	100%	94%	100%	97%	100%	85%	94%	92%	97%	97%	91%	94%	91%
pos occupied <sup>6</sup>	1	1	1	1	1			•••••										4

Table 4D: Analysis of V kappa subgroup 4

											Fra	mew	ork/	11				
amino acid'	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48
Α				32						2					T			
В													•••••••	•				
· c												``					7	
D					•									İ				<u> </u>
Е											1	<u>.</u>		<del></del>				
F																		
G											32					<u></u>		
Н		•				2										÷		
		<u>.</u>															••••	32
K					<u>.</u>				33						32			
<u> </u>			33		; ; ; ;											29	33	
· M		<u>.</u>											: :			,		1
N	33	<u>.</u>	<u>.</u>									••••••	· · · · · · · · · · · · · · · · · · ·					********
Р										31		*********	31	33			••••••	••••••
Q		<u> </u>					32	33				32		······		•	•••••••••••••••••••••••••••••••••••••••	********
R		<u> </u>	<u> </u>				1					1		,	1			•
<u>S</u>		ļ	ļ		•								2					*********
		<u> </u>		1	********		*******	7										
V		<u> </u>	<u></u>					************								4		
W		<u>.</u>			33		*****	•••••••										
X							********		**********					٠				Ţ
Υ		<b>3</b> 3				31												
	ļ							*********	******									
unknown (?)								**********	*********									
not sequenced												,						
sum of seq <sup>2</sup>	33	33	33	33	33	33	33	33	33	33	33	33	33	33	33	33	33	33
oomcaa3	33	33	33	32	33	31	32	33	33	31	32	32	31	33	32	29	33	32
mcaa <sup>4</sup>	N	Υ	L	Α	W	Υ	Q	Q	Κ	Р	G	Q	Р	Р	K	L	L	1
rel. oomcaas	100%	100%	100%	92%	100%	94%	92%	100%	100%	94%	97%	97%	94%	100%	97%	88%	100%	97%
pos occupied <sup>a</sup>	1	1	1	2	1	2	2	:			2		:	•••••••••••••••••••••••••••••••••••••••	2	2	1	2

Table 4D: Analysis of V kappa subgroup 4

2 40. Alialysis 01 V			- 3.	•	CDR	11										•••		
amino acid'	49	20	51	52	53	54	55	56	57	28	59	09	61	62	63	64	65	99
А			30															İ
В								; •••••••• • • • • •		}	(*************************************	• !		•		•		
· C						*******	•	***************************************	• • • • • • • • • • • • • • • • • • •	**************************************	<del></del>	•					••••••	
D										·		33					******	
E							32						***********		•••••		*********	
F				-						••••••	••••••	••••••	********	33	*********	•••••	*********	••••••
G			·						33		***********		*********			33	•	33
Н										·					•	••••••		
l					1			•••••		**********	****		********	•	*******			*********
K													*******		•		*********	*******
L						·												********
М											*********			•••••				••••••
N					2	**********		*********	_	•		*******					***************************************	••••••••
Р				1	•••••			*******		•••••	33	*********	1	••••••	•••••••			
Q		•	************				••••••	•••••		**********	•••••••			••••••	•••••••			
R						33				•••••	**********		32		••••••			
S			1	31	1			33					•••••		32		33	*********
T			2	1	29								•••••••				·i	
V							1			33		***************************************	***************************************	***************************************				
W		33					•••••••••••••••••••••••••••••••••••••••				••••••	***************************************						
X								**********			••••••		************	***************************************				••••••
Υ	33										***************************************		•••••••••••••••••••••••••••••••••••••••			************		
<b>-</b> .																		
unknown (?)								********			•	***************************************		**********			••••••	•••••••
not sequenced								***********					••••••	***************************************		***************************************	•••••••••••••••••••••••••••••••••••••••	******
sum of seq <sup>2</sup>	33	33	33	33	33	33	33	33	33	33	33	33	33	33	33	33	33	33
oomcaa <sup>3</sup>									:		:	•••••••••••••••••••	•••••••••••••••••••••••••••••••••••••••	33	•••••••••••••••••••••••••••••••••••••••	•••••••••••••••••••••••••••••••••••••••	•••••••••	*******
mcaa <sup>4</sup>	Υ	•	Α	S	T	R	Ε	S	G	٧٠	Р	D	R	F	S	G	•••••••••••••••••••••••••••••••••••••••	G
rel. oomcaa'	100%	100%	91%	94%	%88	100%	97%	100%	100%	100%	100%	100%	97%	100%	97%	0001	%00 I	%00I
pos occupied <sup>6</sup>	1	1		:							:	•••••••••••••••••••••••••••••••••••••••	:	1	:			

Table 4D: Analysis of V kappa subgroup 4

ic 40.7 mary 313 01 V	-171		- 3.			ame	wor	k III										• ;
amino acid'	29	89	69	20	71	72	73	74	75	9/	77	78	79	8	81	82	83	84
Α														33			:	32
В														<del></del>				
. C							-			<del></del>				-				
D				32										<u> </u>		33		
E										···········				•	33			
F.					32				-	-	<del></del>			•				
G		33		1	<del></del>					······		<u> </u>	<del></del>	<del></del>				1
Н										······		<u>.</u>	············	<u></u>			••••••••••••••••••••••••••••••••••••••	
1									33	<del></del>				*************	•••••		••••••	
K															••••••	•••••	•	
<u>L</u>							33					32		• • • • • • • • • • • • • • • • • • •	***********			
. M										,		1				<b></b>		
N										2	1		********		•••••			
Р							•	*******			*******	••••••••	*******		••••	•••••	•	
Q				-					-				32				*****	
R								·					1		***************************************		•••••	
S	33		••••••					•		30	32							
T			33			33		33		.1								
V			••••••		1												33	
W			•••••		********													
X			,															
Υ																		
								*********										
unknown (?)																		
not sequenced																••••••		
sum of seq?	33	33	33	33	33	33	33	33	33	33	33	33	33	33	33	33	33	33
oomcaa,	33	33	33	32	32	33	33	33	33	30	32	32	32	33	33	33	33	32
mcaa <sup>4</sup>	S	G	Ţ	D	F	T	L	Ţ	1	S	S	L	Q	Α	Е	D	٧	Α
rel. oomcaa <sup>s</sup>	100%	100%	100%	97%	97%	100%	100%	100%	100%	91%	97%	97%	97%	100%	100%	100%	100%	97%
pos occupied <sup>6</sup>	1	1	1			:	1	1 &	•••••••••••••••••••••••••••••••••••••••		2		<u>-</u>	1	1	1	1	

Table 4D: Analysis of V kappa subgroup 4

			. <u>*                                    </u>						·		C	DR I	111					
amino acid'	82	98	87	88	83	90	91	92	93	94	95	V	8	ပ	۵	ш	щ	96
A										1								
В									*******			•••••				.,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,		
·				33					•••••									
D			******					1	1				•••••					<b></b>
E																		
F ·			1					1										
G									2	_								
. н			1		3									•				
İ										2								
K																		
L						1		2		1	3							1
· M																		
N									4	4								
Р .										1	29	1						4
Q					30	32					1							1
R									1		·	1						2
S							2		23	2								1
T									2	22								
V	33										.,							
W																		2
X																		************
.Υ		33	31				31	29										1
												13	15	15	15	15	15	3
unknown (?)								*********			*********							
not sequenced												18	18	18	18	18	18	18
sum of seq'	33	33	33	33	33	33	33	33	33	33	33	15	15	15	15	15	15	15
oomcaa³	33	33	31	33	30	32	31	29	23	22	29	13	15	15	15	15	15	4
mcaa•	٧	Υ	Υ	С	Q	Q	Υ	Υ	S	T	Р	-	-	-	-	-	~	Р
rel. oomcaas	100%	100%	94%	100%	91%	92%	94%	9/088	20%	67%	9/88	87%	100%	100%	100%	100%	100%	27%
pos occupied <sup>6</sup>	1	1	3	1					6	7	3	3	1	1	1	1	1	8

Table 4D: Analysis of V kappa subgroup 4

violy 5.5 or vinappa						Fr	ame	worl	< IV		<del></del>			1
amino acid'	97	98	66	100	101	102	103	104	105	106	⋖	107	108	sum
Α														183
В		•	[	••••••••••••••••••••••••••••••••••••••		••••••			<b></b>		<b>*</b> ······		<b></b>	
С		************	•											68
D	1	<del></del>						<u> </u>			<u></u>	ļ		154
E			•••••			•	·········		14					105
. F		15		•	************				•••••				••••••	82
G			15	4	15						<del></del>			228
Н			•••••			·			•	••••••	·······	•••••••	•••••	6
									•••••	14	•	•••••		135
K						••••••	14			••••••		13		158
L						••••		4			********	•••••		258
M	1									***********	•••••	*********	********	27
N												1	·	136
P						1								195
Q				11				1			,			264
R							1	·	1			1	11	116
S	2									1				499
T	12					14							411111111	236
V								9						196
W							-	1						69
X														
Y							-							254
-											15			106
unknown (?)														
not sequenced	18	18	18	18	18	18	18	18	18	18	18	18	22	518
sum of seq'	15	15	15	15	15	15	15	15	15	15	15	15	11	
oomcaa³	12	15	15	11	15	14	14	9	14	14	15	13	11	
mcaa <sup>4</sup>	Ţ	F	G	Q	G	T	K	٧	Ε	1	-	Κ	R	-
rel. oomcaa'	80%	100%	100%	73%	100%	93%	93%	%09	93%	93%	100%	87%	%00 I	
pos occupied <sup>e</sup>	3	1	1	2	1	•••••	2	:	2	2	1	3		

Table 5A: Analysis of V lambda subgroup 1

											Fran	new	ork l						
amino acid¹	_	7	က	4	2	9	7	&	6	10	=	12	13	14	15	91	17	18	19
Α						-					19		18	20					
В						·			• • • • • • • • • • • • • • • • • • •					•••••					
· C				• • • • • • • • • • • • • • • • • • •	********							•••••							
D		•••••			•••••		*****					••••••		•••••					
Ε				••••••			••••••	•••••••			······	*********	••••••		••••••			1	
F .	***********				•••••••							•••••••							••••••
G	·			•	•				••••••				22		•••••	42			
Н	2			•••••								•••••			••••••				
			1					••••••			1	******			••••••	••••			
К								•	••••			•••••			• • • • • • • • • •	•••••		14	
L			1	41		•		********		•	1				•••••••		•		
М			•••••			•		••••	•••••				•••••		•••••		•••••		
N				•••••		•••••		*********		********		•••••	•••••			•••••	•••••		,
Р		•				•••••	41	41						1	41				
Q	22		1	•	•••••	41	************		•			•••••					42		
R							***************************************		********		***************************************	*******		٠	•••••••			25	
S		39							41			41			1			1	
Т					41							•		19				1	***************************************
V		1	38								20		1	1	••••••				42
W															*******				
X																			
Y															•				
Z	16																		
_										41									
unknown (?)																			
not sequenced	2	2	1	1	1	1	1	1	1	1	1	1	1	1					
sum of seq²	40	40	41	41	41	41	41	41	41	41	41	41	41	41	42	42	42	42	42
oomcaa,	22	39	38	41	41	41	41	41	41	41	20	41	22	20	41	42	42	25	42
mcaa•	Q	S	٧	L	Ţ	Q	Р	Р	S	_	٧	S	G	Α	Р	G	Q	R	٧
rel. oomcaas	55%	%86	93%	100%	100%	100%	100%	100%	100%	100%	49%	100%	54%	49%	%86	100%	100%	%09	100%
pos occupied <sup>e</sup>					1	1	1	1	1	1	4	1	:	•		•••••			1

e SA: Affaiysis of											CE	RI							-
amino acid'	70	21	22	23	24	25	56	27	۵	ш	28	29	30	31	∢	32	33	34	35
Α	2			,				1				2	2			1			
В																			
C ·				42															••••••
D										3			3	1		3		1	
E													1						
F					1				1						1	1			••••••
G						42	3	1			2	39	4	2					.,,,,,
Н														2		2		2	
1	1	41								1	37							1	
K							1			1			1						
L		1									1								
М										-	1								
N								2	1	37			13	31	2		1	9	
Р																1			
Q																1			
R							1	1					5			į			
S	1		42		38		34	34	38				13	1	1	3		19	
T	38	·			3		4	3	2			1		1		7		2	
V					4						1					2	40		********
W																			42
X																			
Y														4	1	20		7	
Z																			
-										·					36				
unknown (?)																			
not sequenced														·	1	1	1	1	
sum of seq <sup>2</sup>	42	42	42	42	42	42	42	42	42	42	42	42	42	42	41	41	41	41	42
oomcaa3	38	41	42	42	38	42	34	34	38	37	37	39	13	31	36	20	40	19	42
mcaa <sup>4</sup>	T	İ	S	С	S	G	S	S	S	N	1	G	N	N	-	Υ	٧	S	W
rel. oomcaas	<b>%06</b>	%86	100%	100%	<b>%</b> 06	100%	81%	81%	%06	. %88	98%	93%	31%	74%	%88	49%	98%	46%	100%
pos occupied <sup>6</sup>		•	†···	: :	Ţ <del></del>	<del>}</del>	<del>;</del>	•	:	:	:						. 2		

Table 5A: Analysis of V lambda subgroup 1

				•		Fran	iewo	ork II											
amino acid'	36	37	38	39	40	41	42	43	44	45	46	47	48	49	20	51	52	53	54
А							4	40									1		
В												<u></u>							
C																			
D						1									13	10	8		
E										2					5			1	
F	1			4										1					
G						39									1		٠		
Н	1	1	6	1						·				1				1	
1												<u> </u>	40		1				
K							1			35					1	1		18	
L			1	31							41	40						1	1
М							1						1					1	
N										1					3	28	30	2	
Р					42	1			42			į							
O .		39	34															15	
R		2		1		1				4		<u></u>			7			2	40
S								1							9	2	3	1	
T							36	1							1				
V			1	5		-					1	2	1		•••••				
W		,		)										•••••	•••••				1
X									********										
Υ	40													40	1	1			
Z																			
-																			
unknown (?)																			
not sequenced				<u> </u>	<u> </u>												<u> </u>		
sum of seq <sup>2</sup>	42	42	42	42	42	42	42	42	42	42	42	42	42	42	42	42	42	42	42
oomcaa,	40	39	34	31	42	39	36	40	42	35	41	40	40	40	13	28	30	18	40
mcaa*	Υ	Q	Q	L	Р	G	Ţ	Α	Р	K	L	L	ı	Υ	D	N	N	K	R
rel. oomcaa <sup>s</sup>	95%	93%	81%	74%	100%	93%	%98	95%	100%	83%	%86	95%	95%	95%	31%	%29	71%	43%	95%
pos occupied	•		:	Ŧ	1	-		•	-	-	-	:		:	10		<u> </u>	•	;

	CD	RII			·									•					
amino acid'	55	26	٧	8	ပ	۵	ш	57	58	59	9	61	62	63	64	65	99	<b>4</b>	В
Α	1														5				
В		<u></u>	<u></u>																
· C	<b>.</b>																		
D			<u></u>								38								
E	ļ																		
F.													38						
G								41			2				36				
Н											1								
									17				3	**************************************				•••••	
К								***********					*********	: :	••••••	•••••	38		*******
L		1						•		1			••••••			••••			******
M							••••••	••••••					••••	***************************************	•••••	••••			*******
N				**********	*********	********		•••••								*******			
Р	38							********		38			• • • • • • • • • • • • • • • • • • • •	••••••					*********
Q									•				•						••••••
R									••••			42	•••••••	••••••		•••••	4		
·S	2	40								2				42		42			*********
T															1	********			
V									24				1			********			
W													•	**********	•	********			********
X														*******		***********			•••••••
Y														-		••••••		•••••••••••••••••••••••••••••••••••••••	•
Z													••••••••	********		**********		•••••••••••••••••••••••••••••••••••••••	
-			41	41	41	41	42											42	42
unknown (?)		4									***********								
not sequenced	1	1						1	1	1	1								
sum of seq²	41	41	41	41	41	41	42	41	41	41	41	42	42	42	42	42	42	42	42
oomcaa³	38	40	41	41	41	41	42	41	24	38	38	42	38	42	36	42	38	42	42
mcaa'	Р	S	-	-	-	-	-	G	٧	Р	D	R	F	S	G	S	Κ	-	-
rel. oomcaa <sup>s</sup>	93%	%86	100%	100%	100%	100%	100%	100%	29%	93%	93%	100%	%06	100%	96%	0001	<b>%06</b>	100%	100%
pos occupied <sup>6</sup>		2	1	1	1		1		2		;		:	•••••••••••••••••••••••••••••••••••••••	•••••••••••••••••••••••••••••••••••••••	1		•••••••••••••••••••••••••••••••••••••••	

Table 5A: Analysis of V lambda subgroup 1

				Fra	ame	work	: III												
amino acid'	29	89	69	70	71	72	73	74	75	9/	77	78	79	8	81	82	83	84	85
A		1	3		41			24						2				38	1
В																			
· c																			
D		1													1	41			37
E													1	••••••	24		42		1
F.									*********		••••	*********	•			•••••			*******
G		40						17		1	42	*********		•••••	15				**********
Н													1						2
									41						•••••				1
K									***************************************			•			•••••				*********
L							42	••••••		***********		41							*********
М			************						•••••										*******
N				**********												1			*********
Р									•••••••					2					*********
Q				•					**********			•••••	31					••••••	*******
R						***************************************	***********		•••••••••••••••••••••••••••••••••••••••				8					•••••••	
S	42		1	42		24			***************************************	20				20				1	
Т			38			18			••••••••	21				17				3	
V				-,	1			1	1			1		1					******
W						**********							1		2				
X																			*****
Y																			
Z																			
-																			
unknown (?)				-															*******
not sequenced																*********			•••••
sum of seq <sup>2</sup>	42	42	42	42	42	42	42	42	42	42	42	42	42	42	42	42	42	42	42
oomcaa <sup>1</sup>	42	40	38	42	41	24	42	24	41	21	42	41	31	20	24	41	42	38	37
mcaa'	S	G	T	S	Α	S	L	Α	١	Ţ	G	L	Q	S	Ε	D	Ε	Α	D
rel. oomcaas	100%	95%	%06	100%	%86	57%	100%	57%	%86		100%	%86	74%	48%	57%	%86	100%	%06	88%
pos occupied <sup>6</sup>	1		3	•••••••••••••••••••••••••••••••••••••••			1			3	•••••••••••••••••••••••••••••••••••••••	••••••	:	:		:	•••••••••••••••••••••••••••••••••••••••	•••••	<u></u> 5

## WO 97/08320 Table 5A: Analysis of V lambda subgroup 1

(C 5/1. / mary5/5 5									•	CD	D III	<u> </u>					•		
			i			•			<u>.</u>		R III							<u> </u>	
amino acid¹	98	8	8	88	6	6	92	6	96	95	<b>∀</b>	മ	ပ —		ш	<u> </u>	6	65	<u>8</u>
Α				22	15			1				16					4	1	
В								**********						,,,,,,,,,,,					
С			42											*******					
D							39	17			7								
E				į								1					1		
F		2								1	<b>400000</b>								36
G				14				1				-17	1				5	1	
Н		.1				İ							1						
1					<u></u>						1							1	
κ				į		<u>i</u>					1								
L				1						37	*****		1					1	
М																		1	
N							2	2			9	1							
Р				į		<u></u>				1							6		
Q				3															
R									5	1	2						2		
S					4			17	35		18		1			•••••	1		
T					22			1	1		1		-						
V				1				1		1		2					9	34	
W						38											7		
X																			
Υ	42	39				3		1								•••••	3		
Z								-											
-											2	4	35	39	38	38	1		
unknown (?)																			
not sequenced				1	1	1	1	1	1	1	1	1	3	3	3	3	3	3	4
sum of seq <sup>2</sup>	42	42	42	41	41	41	41	41	41	41	41	41	39	39	38	38	39	39	36
oomcaa,	42	39	42	22	22	38	39	17	35	37	18	17	35	39	38	38	9	34	36
mcaa <sup>4</sup>	Υ	Υ	С	Α	T	W	D	D	S	L	S	G	-	-	-	-	٧	٧	F
rel. oomcaas	100%	93%	100%	54%	54%	93%	95%	41%	85%	<b>%</b> 06	44%	41%	<b>%06</b>	100%	100%	100%	23%	87%	100%
pos occupied	1	i								•	:			1	1	1	10		

Table 5A: Analysis of V lambda subgroup 1

•			I	ram	ewo	rk IV	,	-				
amino acid'	66	901	101	102	103	104	105	106	⋖	107	108	sum
Α												285
В												
С												84
D												224
E		1					į					81
F							į	į		į		87
· G	36	31	36							26		559
Н										<u></u>		25
1					<u> </u>		<u></u>					188
K					30							141
L						25			34			344
М										• • •		5
N					1							176
Р											1	296
Q					3				1		18	251
R					1			į		2		156
S		1								2		720
T		3	*********	-36	1		36			<u></u>		359
V						11		36	1			282
W										1		92
X												
·Υ												202
Z			-								٠	16
-												524
unknown (?)						•••••					*********	
not sequenced	4	6	6	6	6	6	6	6	6	10	22	141
sum of seq'	36	36	36	36	36	36	36	36	36	31	19	
oomcaa,	36	31	36	36	30	25	36	36	34	26	18	
mcaa*	G	G	G	T	K	L	T	٧	L	G	Q	
rel. oomcaas	100%	%98	100%	100%	83%	%69	100%	100%	94%	84%	95%	
pos occupied	1	4	1	1	5	2	1	1	-3	4	2	

27

											Fra	mew	ork/	I		-			
amino acid'	-	7	က	4	S	9	7	œ	6	2	=	12	13	14	15	16	17	81	19
Α			35					30			6		1	1		Ī			
В				:						<u> </u>				<u> </u>	<u> </u>		<u> </u>	<u> </u>	
· c				-							<u></u>		<u> </u>	<del></del>	<u> </u>	<u> </u>	<u> </u>	<u> </u>	<u> </u>
D			·			<u> </u>		<u> </u>			<del></del>		<del></del> -		<u>†</u>	1	<b></b>	<del></del>	
E			•							<del></del>	<del></del>		<del> </del>		<del></del>		<u> </u>	<del>!</del>	<del></del>
F .		*******	•	*********					••••••••••••••••••••••••••••••••••••••			********	<del> </del>	<del></del>	<del>•</del>	<u> </u>	<u> </u>	<del></del>	<b></b>
G		·		·····			ļ		}		••••••	••••••	42	<del> </del>	<del></del>	42	······	<b></b>	ļ
Н	2			•			······					••••••			<del></del>		1		ļ
1			1									•••••	······		<u></u>		•••••		28
K									*******						<u></u>				
L				40			• • • • • • • • • • • • • • • • • • •		•••••			•••••			3			•••••	1
М			<u> </u>					•••••	********				*********						
N			<u> </u>	••••••					••••••			•••••	•••••						••••••
P		•••••••	<u> </u>		**********	••••	42	6	*******	••••••			••••••	•••••	40				*********
Q	22	********	4	••••••	**********	41	•••••	• • • • • • • • • • • • • • • • • • • •	•••••••	•••••	•		********	*********			42		******
R							•••••	6	1			*******	•						
S		41							40			42		42				43	,
T				********	42		********		1				******				•••••••••••••••••••••••••••••••••••••••	*********	,
V		1	2	*********	***********	•••••	•••••				36	***************************************	***********		*********				14
W		****		*******	•••••••••••••••••••••••••••••••••••••••				*********										
Χ		********										*******	********			••••••			
Υ							•••••••		•••••			••••••				•••••••••••••••••••••••••••••••••••••••	•••••		
Z	16								••••••								•••••••••••••••••••••••••••••••••••••••		
-										42									=
unknown (?)						1			••••••	İ			•••••••••••••••••••••••••••••••••••••••				••••••		
not sequenced	3	1	1	3	1	1	1	1	1	1	1	1	*******	·····			•••••	••••	
sum of seq <sup>2</sup>	40	42	42	40	42	42	42	42	42	42	42	42	43	43	43	43	43	43	43
oomcaa³	:		:		·······	•••••••••••••••••••••••••••••••••••••••	••••••••	•••••••••••••••••••••••••••••••••••••••	•••••••••••••••••••••••••••••••••••••••		36		•••••••••••••••••••••••••••••••••••••••	•••••••••••••••••••••••••••••••••••••••	••••••	42	•••••••••••••••••••••••••••••••••••••••		•••••••••••••••••••••••••••••••••••••••
mcaa¹	Q		Α	L	T	Q	Р	Α	S	-	٧	S	G	S	Р		Q	S	1
rel. oomcaa <sup>s</sup>	55%	%86	83%	100%	100%	%86	100%	71%	95%	100%	%98	100%	98%	%86	93%	%86	%86	100%	965%
pos occupied <sup>6</sup>		:	•	1		1				••••••	•	1	2	2	2		2	······	3

Table 5B: Analysis of V lambda subgroup 2

											CI	RI							
amino acid'	20	21	22	23	24	25	76	27	۵	ш	28	29	30	31	⋖	32	33	34	35
Α					3		1						1			1			
В								••••	••••										
· C		**********		42					1					1					
D		•••••						•••••		39		1	4		5				
E								•••••	•••••						1				
F.		1											1			4			
G		***********				43		1				39	26						
Н								1							1	1			
1		41			1						6								
. K															4				
L		1														4			
М																			
N								1	3	4		1	4	3	28				
Р								1											
Q																			*********
R									1				2						********
S			42		3		3	35	38				5	1	2	4	1	42	
T	43				36		39	3				1		1					
V											37						41		
W												,							43
X																			
Υ								1				1		37		29			
Z																			
_	,														1				
unknown (?)															1				
not sequenced			1	1													1	1	
sum of seq <sup>2</sup>	43	43	42	42	43	43	43	43	43	43	43	43	43	43	43	43	42	42	43
oomcaa³	43	41	42	42	36	43	39	35	38	39	37	39	26	37	28	29	41	42	43
mcaa'	T	١	S	С	T	G	T	S	S	D	٧	G	G	Υ	N	Υ	٧	S	W
rel. oomcaas	100%	95%	100%	100%	84%	100%	91%	81%	98%	91%	%98	91%	%09	. %98	65%	67%	%86	%001	100%
pos occupied	:	:	:	1				:			:					:			- 1

Table 5B: Analysis of V lambda subgroup 2

						Fran	new	ork II											
amino acid'	36	37	38	39	40	41	42	43	44	45	46	47	48	49	20	51	52	53	54
Α					1	- 4		40											
В																			
С																			
D				1		2									20	1	2	1	
E															20			2	
F.	2													7		1			
G						36			*******	•				********	2	2		1	*********
Н			2	34			*******		******		•				•••••		*****	1	
1							1				1	9	43		••••••		1		
K						**********	40		******	41				,			1	21	*********
L ·			1	1					••••••		38	6		*******					10000000
М		·							***************************************			26	4				1		********
N				2	·									**********	1	••••••	8	12	********
Р	·				41				43										100 00000 84
Q		41	39							2									
R		1			·		1		•••••					•			2		43
S					1		•						***************************************	2			21	3	*********
T						***************************************	1		*******		••••••	İ		•••••••••••••••••••••••••••••••••••••••			7		
V .						1		3	*********		4	2				39	······		***********
W							,							•••••••					*********
Χ									~			<u>-</u>		••••••					*******
Υ	41			5			•		•••••••••		•	••••••		34				2	*******
Z									••••••	•			••••••		•		••••••		*********
<u>.</u>						į													
unknown (?)		1	1								••••••		•••••••••••••••••••••••••••••••••••••••	•••••••••••••••••••••••••••••••••••••••	**********				*******
not sequenced																			
sum of seq <sup>7</sup>	43	43	43	43	43	43	43	43	43	43	43	43	43	43	43	43	43	43	43
oomcaa¹	41	41	39	34	41	36	40	40	43	41	38	26	43	34	20	39	21	21	43
mcaa <sup>4</sup>	Υ		Q			:	:	Α	:	:		М		:	•••••••••••••••••••••••••••••••••••••••	٧	•••••••••••••••••••••••••••••••••••••••	:	********
rel. oomcaas	95%	95%	91%	79%	95%	84%	93%	93%	100%	95%	%88	%09	100%	29%	47%	91%	49%	49%	100%
pos occupied <sup>6</sup>	•				:	:		2	:				:		4	•	8	:	********

Table 5B: Analysis of V lambda subgroup 2

	CD	R II							<del></del>										
amino acid'	55	26	⋖	മ	ပ	۵	ш	22	28	59	09	61	62	63	64	65	99	4	æ
Α															2				
В																			
С																1			
D											17		.,						
E																			
F													42						
G								43	1						41				
Н											2								
1									3										
К												<u> </u>					42		
L											1		1				· [		
М																			
N											19								
Р	43									15									
Q																			
R												43					1		
S		43								28	2	<u>į</u>		43		42			
T																			
V									39										
W																			
X																			
Y				••••••	•••••						2								
Z																			
		•••••	43	43	43	43	43											43	43
unknown (?)														*********		•••••			
not sequenced	•															-			
	<u> </u>									:		43			**********	**********			*********
oomcaa,	43		43	43	43	43	43	43	39		19	43	*******	43	41	42	42	43	43
mcaa*	Р	S	-	_	-	-	-	G	٧	S	N	R	F	S	G	S	K	-	-
rel. oomcaa <sup>s</sup>	100%	100%	100%	100%	100%	100%	100%	100%	91%	65%	44%	100%	%86	100%	95%	%86	98%	100%	100%
pos occupied <sup>6</sup>	1	1	1	1	1	1	1	1	3	2	6	1	2	1	2	2	2	1	1

Table 5B: Analysis of V lambda subgroup 2

				Fra	ame	work	c III	٠.								•			·
amino acid¹	29	89	69	. 02	71	72	73	74	75	9/	77	78	79	80	81	82	83	84	85
Α		3		1	43									36				43	
В			<u>.</u>	<u></u>															
. С	<u>.</u>																		
D	·	1	2												3	42			39
E			: : :								1				38		43		
F .						••••													
G		39									42				1				
Н																			2
<u> </u>		•							35										
K			1			••••													
L		•				*******	43					43							
M						******													
N			38			**********					٠				1	1			1
Р						P = P = 0 = 0 = 0	•	*******						2					
Q			********										41						
R													· 2						
S	42			1	į	43				42				į					
T			1	41			V PE PRO 0000	43		1				2					
V							••••••		8					3				· ·	
W							•	********											
X								********											
Y								•••••											
Z								~											
								*******											
unknown (?)			1																1
not sequenced	1																		
sum of seq <sup>2</sup>	42	43	43	43	43	43	43	43	43	43	43	43	43	43	43	43	43	43	43
oomcaa <sup>3</sup>	42	39	38	41	43	43	43	43	35	42	42	43	41	36	38	42	43	43	39
mcaa•	S	G	N	Ţ	Α	S	L	T	ı	S	G	L	Q	Α	Ε	D	Ε	Α	D
rel. oomcaa <sup>5</sup>	100%	91%	98%	%S6	100%	100%	100%	100%	81%	%86	98%	100%	95%	84%	88%	%86	100%	100%	91%
pos occupied <sup>6</sup>	1	3		:	1	1	1			:		1		:	4		1	1	

Table 5B: Analysis of V lambda subgroup 2

										CD	R III								<del></del>
amino acid'	98	87	88	83	90	91	92	93	94	95	∢	8	U	۵	ш	ட	96	97	86
Α				2	1		21		1								1	1	
В											••••••		*********	***********	• • • • • • • • • • • • • • • • • • •				
· c			43	11															
D								3	1	2							1		
Е							1	1											
F .		3				3				1		1					5		42
G							1	21	3	4							1		
н						1													
I							1	1		1	2						1	7	
К										3									
L												1	1				6	5	
М																	1	1	
N									.5	7	5						1		
Р								1				4							
Q						***********				1	2								
R							2		3			1					5		
S·		1		30	41			12	23	14	9						1		
T							16	4	4	3	21								
V							1										11	28	
W																	5		<u></u> j
X																			
Y	43	39	-			39			1	6							4	************	
Z																			
										1	3	36	42	43	43	43			
unknown (?)									2					********					<u></u>
not sequenced					1						1							1	_1
sum of seq'	43	43	43	43	42	43	43	43	43	43	42	43	43	43	43	43	43	42	42
oomcaa,	43	39	43	30	41	39	21	21	23	14	21	36	42	43	43	43	11	28	42
mcaa'	Υ		С	S	S	Υ	Α	G	S	S	Τ	-	-	-	-	-	٧	٧	F
rel. oomcaa <sup>s</sup>	100%	91%	100%	20%	%86	91%	49%	49%	53%	33%	20%	84%	%86	100%	100%	100%	26%	67%	100%
pos occupied <sup>6</sup>	:	:												**********		1	13		

				Fra	mew	ork	IV					7
amino acid'	66	9	101	102	103	104	105	106	4	107	108	sum
Α		1				Ī			I			280
В			<u> </u>			<u> </u>			<u> </u>		<del></del>	
С		<u> </u>	-	<u> </u>	**********	<del></del> -		· -	·	<del></del> -	<u> </u>	99
D		-		<del></del>	<u> </u>	· •	·	· • • • • • • • • • • • • • • • • • • •	<del></del>	·	<del> </del>	188
E		1	<u> </u>			<u> </u>	· · · · · · · · · · · · · · · · · · ·	·	<u> </u>		<del> </del>	107
F			·	<u> </u>	·	<u> </u>	·	<u>†</u>	·	·	<del></del>	113
G	42	33	42	•		<b>†</b>	·	·	<b></b>	19	·	567
Н	·	•	·			•			·		<u></u>	48
	1	<u></u>			·	<del> </del>	1	·•	<u> </u>		<u> </u>	184
K		<u></u>	·	<u></u>	36	<u></u>	<u> </u>	<u> </u>	<u> </u>		<u></u>	189
L				<u></u>		28	<u></u>	<u> </u>	40		<u></u>	264
M	1						<del>.</del>	<u> </u>				29
N			<b></b>		1	<u></u>	<u></u>		<u></u>			146
P	·	**********				<b>÷</b>	ļ	<u> </u>				238
Q		••••••		********	1	<b>†</b>	!	<u> </u>	<u>.</u>		14	1
R		1		********	2	ļ	·	<del></del>	••••••••••••••••••••••••••••••••••••••	4		121
S		*********				ļ	1	<u></u> .		2		831
T		7		41	<b></b>	<u> </u>	40	 !				398
V		*********		•••••••		14		42	1			327
W		•••••		********		•••••	•••••					48
X				********	••••••	••••••	********		•••••			.0
Υ		*********		********	1	••••••	•••••		********			285
Z		********		••••••	•				•••••			16
-											╡	555
unknown (?)			i					••••••				8
not sequenced	1	1	1	2	2	1	1	1	2	15	28	80
sum of seq²	42	42	42	41	41	42		==		25	14	
oomcaa³			42	•••••••••••••••••••••••••••••••••••••••	••••••••••••	••••••••		•••••••••••••••••••••••••••••••••••••••	40	19	14	
mcaa⁴	G	G	G	T	Κ	L	Т	٧	L	G	Q	
rel. oomcaas	%00I	%62	%00I	%00 I	988%	9/2/9	95%	%00I	%86	76%	<b>%00</b>	
000 000-1-46			····		······································	••••••	:			•••••••••••••••••••••••••••••••••••••••	5	
pos occupied <sup>6</sup>	1	4	1	1	5	2	3	1	2	3	1	

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Table 5C: Analysis of V lambda subgroup 3

į											Frar	new	ork l						
amino acid'	_	2	က	4	2	9	7	œ	6	10	=	12	13	14	15	16	17	18	19
А					1		1	2	7					20	1				27
В		•••••						*******	**********										
. C				•					**********										
D		*********	5	********			10		**********										******
E	.,		20										1			1			
F .	1	1										1			1				
G			1													37			
Н																			•
l																			•
. К																	2	••••••	
L				37							4		1		9				********
М				********		•••••••								•••••••					
N	***************************************	***************************************		********	********				•				**********						*******
Р				••••••	••••••		26	35	1		•••••				27			•••••••	1
Q	4	*******	4	********		38			•••••••		•••••••••••••••••••••••••••••••••••••••		***************************************		•••••••		36		•••••
R		***********		*******		•					***************************************		•••••••						
S	13	14			1		1		28			37	********	18					
Т		•		•	36			1	•••••••••••••••••••••••••••••••••••••••					••••••				38	
V			8	1					2		34		36	<u>-</u>					10
w				••••					••••••				•			*********		**********	
х		••••••	••••	****		•							••••••	*********					
Y		23		•		•	•••••	••••••											
Z		••••••		•		•													
-	20			_						38									
unknown (?)		••••••	********		••••••	••••••		•	***********		•••••					•••••	•••••		
not sequenced				*******	4	*********	••••	**********			*******	••••					,		
sum of seq <sup>2</sup>	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38
oomcaa³				******					***********	***************************************			*********	20	••••••		**********	***************************************	
mcaa'		Υ		L			Р	**********		_	•••••		٧			G		_	Α
rel. oomcaa⁵	%	%		•••••	•••••			••••••		_	•••••		•••••	**********		•••••			•••••
•	53%	•	53%				:	:	74%				95%	53%	71%	97%	92%	<u>5</u>	71%
pos occupied <sup>6</sup>	4	3	5	2	3	1	4	3	4	1				2	4	2	2	1	3

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Table 5C: Analysis of V lambda subgroup 3

ie 5C: Analysis o					, , , , , , , , , , , , , , , , , , ,							DRI							
	_		~		<u> </u>		(0	_		•									<u></u>
amino acid'	~	7	<u>~~</u>	23	7	; ;	5	27	۵	<u>ш</u>	<u>~</u>	- 56	∺ —	'n	<b>▼</b>	<u>~~</u>	<u>~~</u>	34	<u> </u>
Α	ļ	<u>.</u>	1					5		ļ	<u>.</u>	<u></u>	1	1	<u> </u>	ļ <del></del>	21	3	
. В	<b>.</b>	<b>!</b>	ļ							<u>.</u>	<u>.</u>	<u> </u>	<u> </u>	<u>.</u>	<u> </u>	<u> </u>	<u>.</u>		
. С	<b></b>	ļ		38	*****					<u>.</u>	<u> </u>		<u> </u>	<u></u>	<u> </u>	<u> </u>		5	
D							30	1					10			3		1	
E			·				2	2	*********			1	3	6					
F		)			*********									1		2			
G					9	38		1				23	4				٠		
Н							1									2		9	
1		38									9			1					•
K								7					2	13					••••••
L											28								
M.	1													1					
N			2				4	9			1		2			1		2	
Р			1									3							
Q					10									4					
R	25							2				10	1				1		********
S	9		1		19			10					11	2		8		14	
Т	3		33					1				1	4		·				••••
V															•••••	1	15		********
W																			38
· x																			******
Y							1							8	•	20	1	4	
Z														***************************************	•				********
-		į							38	38					37				
unknown (?)													********						•••••••
not sequenced															1	1			•
sum of seq'	38	38	38	38	38	38	38	38	38	38	38	38	38	37	37	37	38	38	38
oomcaa,	:	•		38	:		:				•••••••••••••••••••••••••••••••••••••••		•	*******		*******			38
mcaa¹	R		T	С	S	G	D	S	-	-	L	G	S	K	-	Υ			W
rel. oomcaa <sup>s</sup>	%	<b>‰001</b>	%	100%	9,	100%	ی	2	%	%	 %		<u>چ</u>	Q.	%	Q.	·····-	·····	
	%99	ŏ	87%	•••••	20%	ĭ	79%	26%	100%	100%	74%	61%	29%	35%	100%	54%	55%	37%	100%
pos occupied <sup>a</sup>	4	1	5	1	3	1	5	9	1	1	3	5	9	9	1	7	4	7	1

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Table 5C: Analysis of V lambda subgroup 3

						Fram	ewo	rk II											
amino acidi	36	37	38	33	40	41	42	43	44	45	46	47	48	49	20	51	52	53	54
Α								23								1		1	<u></u>
В					<u></u>		<u> </u>											********	
С					<u> </u>		<u> </u>			į		<u> </u>							
D												į			9	22	2	8	
E			1												5	3		3	
F	3											İ		2			1		
G						36									9	2			
Н					į		1		į	<u></u> j		<u></u>		1	3			1	
1					<u> </u>		į		į	1		<u></u>	28				1		
K				32						İ		<u> </u>		į	2	6	1	13	
L			2			į				6	<b>3</b> 3	1		į					
М	,										1	į	1						
N ·							į			ĺ						1	19	9	
Р					36		1		38			*********							
Q		37	35	1			36								9			1	
R		1		4		2									1	1		1	3
S				1	2			14			į	İ		İ			10	1	
T												į		<u>i</u>		2	4		
٧		·						1		31	4	37	9						
W																		******	
X												į							ļ
Y	<b>3</b> 5													35			.,,		ļ
Z													į						<u> </u>
-																			
unknown (?)																			
ot sequenced		!																	
sum of seq <sup>2</sup>	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	3
oomcaa³	35	37	35	32	36	36	36	23	38	31	33	37	28	35	9	22	19	13	3
mcaa*	Υ	Q	Q	K	Р	G	Q	Α	Р	٧	L	٧	1	Υ	D	D	N	K	1
rel. oomcaas	92%	92%	92%	84%	5%	5%	95%	61%	100%	82%	87%	92%	74%	92%	24%	28%	20%	34%	
pos occupied <sup>a</sup>	•	:	:	:	:	:	:									<u>2</u> 8		<u>က</u> 9	-

Table 5C: Analysis of V lambda subgroup 3

	CI	OR II																	
amino acid'	55	56	V	8	ပ	0	ш	57	28	29	9	61	62	63	64	65	99	∢	8
Α		1											Ī						
В				Ī					<u> </u>		·	<u> </u>	•	<u> </u>	<u> </u>	1	<del></del>	İ	
С	1									· · · · · · · · · · · · · · · · · · ·	<u> </u>		Ì	<del></del>	i	-	<u> </u>	<del> </del>	
D						<u>.</u>		Ī		<del></del>	9		<u> </u>	-	÷	<del> </del>	İ	<del> </del>	
E						-		-			27			<u> </u>	•	-	<del></del>	<u> </u>	
F										•	*******		38		<u> </u>			<del></del>	
G						•		38	<del></del>	<del></del>	<u> </u>		•	ļ	38	·	<del></del>	·····	
Н									]	<u> </u>			<u> </u>		•	ļ		<u></u>	i
· I									37		<u> </u>		•	}		ļ !	<del></del>	<del> </del>	
K						•					• • • • • • • • • • • • • • • • • • •	# # # # # # # # # # # # # # # # # # #	Ī		<u> </u>	<del> </del>	<del></del>	······	
L												<del>***</del>					<u> </u>		
М														***********	<u> </u>				
N									•••••••					********	<u> </u>		21		
Р	37	1							*********	36		*********		•					
Q														**********		•••••			
R												38							
S	1	36			,					1				38		38	12		
Т			·				-							••••••			5		
V														*******		*******		•••	
W												**********		*******					
X																**********	••••••	••••••	*******
Y																		***********	
Z															_		-		*******
_			38	38	38	38	38											38	38
unknown (?)					<u></u>						1					•••••••••••••••••••••••••••••••••••••••		•••••••••••••••••••••••••••••••••••••••	
not sequenced									1	1	1				•••••			•••••	******
sum of seq <sup>2</sup>	38	38	38	38	38	38	38	38	37	37	37	38	38	38	38	38	38	38	38
oomcaa <sup>3</sup>	37	•	•						•			•	38				~~~~~	********	******
mcaa¹	Р	•	-	-	-		-	G	1	Р	Ε	R	F	S	G	S	N	-	-
rel. oomcaas	97%	95%	100%	100%	100%	100%	100%	100%	100%	97%	73%	00%	100%	%00 <sub>1</sub>	<b>%00</b> 1	100%	55%	%001	100%
pos occupied <sup>6</sup>			•	:	:	:		1	:				1	•		••••••	••••••	···	

Table 5C: Analysis of V lambda subgroup 3

				Fra	amev	vork	111										_		
amino acid'	67	89	69	20	71	72	73	74	75	9/	77	78	79	8	81	82	83	84	.85
Α				1	36	1		1				11	1	34				38	
В																			
· C																			
D											·					38			37
E													10		14		38		1
F .														·					
G		37									28				10				
H.			1																
l						1		1	37	1		·			1				
K			1																
L							38								_ 2				
M															10				
N			28		į					1									
Р																			
Q		1											25						
R					·					1	10		1				i		
S	37		2			11				23				1		į			
T	1		6	37	į	25		36		12	į	13		2		į	<u>i</u>		
V					2				1			14	1	1	1	i		į	
W						************										į			
X																			
Y								******											
Z					-														
_						***********										j			
unknown (?)																			
not sequenced																			
sum of seq <sup>2</sup>	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38
oomcaa <sub>3</sub>	37	37	28	37	36	25	38	36	37	23	28	14	25	34	14	38	38	38	37
mcaa <sup>4</sup>	S	G	N	T	Α	T	L	Ţ	1	S	G	٧	0	Α	Ε	D	Е	Α	D
rel. oomcaa <sup>s</sup>	97%	97%	74%	92%	95%	%99	100%	95%	97%	61%	74%	37%	<b>%99</b>	%68	37%	100%	100%	100%	97%
pos occupied <sup>6</sup>											:	:					1	1	2

Table 5C: Analysis of V lambda subgroup 3

•					ОПР				-									_	
				<u> </u>		<del>-</del>					R III		-			····		<u> </u>	· <u>-</u>
amino acid'	98	87	88	8	90	91	92	93	94	95	Α	80	ပ	۵	ш	·	96	97	98
Α					13	3	2			1	2					Ī	4		
В		<u>.</u>																	
· c			38												:				
D		<u></u>	<u>.</u>				32	1	1		6								
E	<b>.</b>	ļ 	ļ	1			400040000					2					2		
F	<b></b>	2						2											35
G	<u> </u>								3	14	3			1			3	1	
Н												12	1						
1																		4	•
K											1							000,000000	
<u> </u>				1				1		1		1	1		·		4	2	•
M				********					1								1	1	
N				10			2	1	2		10	1			•••••				
Р				*******					1	*********			3				1		
Q				25		<u></u>				1	1								
R						10		1	2			2							
S				1	14	1		28	26	13		1				1			
T						1		3		7	2								
V					11												18	28	
W						23	•										1		
X																			
Y	38	36				<u> </u>	1		1		1	3	1				3		
Z																			
											10	15	31	36	37	36		1	
unknown (?)																			
not sequenced							1	1	1	1	2	1	1	1	1	1	1	1	3
sum of seq <sup>2</sup>	38	38	38	38	38	38	37	37	37	37	36	37	37	37	37	37	37	37	35
oomcaa¹	38	36	38	25	14	23	32	28	26	14	10	15	31	36	37	36	18	28	35
mcaa'	Υ	Υ	С	Q	S	W	D	S	S	G	N	-	_	-	_	-	٧	٧	F
rel. oomcaas	100%	95%	100%	%99	37%	61%	%98	<u> 76%</u>	20%	38%	28%	41%	34%	92%	%00 I	97%	%61	76%	%00 I
pos occupied <sup>6</sup>	1	2	1	5	3	5	4	7	8	6	9	8	5	2	1	2	:	••••••	

Table 5C: Analysis of V lambda subgroup 3

				Fram	ewo	rk (\	/					
amino acid'	66	100	101	102	103	104	105	106	∢	107	108	sum
Α												265
В		••••••					•••••					
С		•	•							1		82
D												225
E					2							145
F											**********	90
G	35	31	35						••••	24		461
Н					•••••							32
ı												160
κ					30							110
L		••••••				28			33			233
М									-			17
N		******				•••••••••••••••••••••••••••••••••••••••	•••••			••••••	*******	126
Р					*******	•••••••••••••••••••••••••••••••••••••••			1		*******	249
Q											7	275
R					2							154
S										2		501
Т		4		35			35					347
V			·			7		35				308
w												62
X												
Υ												211
Z												
												603
unknown (?)	<u>.</u>											1
not sequenced	3	3	3	3	4	3	3	3	4	11	28	89
sum of seq²	35	35	35	35	34	35	35	35	34	27	7	
oomcaa <sub>3</sub>	35	31	35	35	30	28	35	35	33	24	7	
mcaa*	G	G	G	T	K	L	T	٧	L	G	Q	
rel. oomcaa'	100%	%68	100%	100%	88%	%08	100%	100%	92%	%68	100%	
pos occupied <sup>a</sup>	1	2	1	1	3	2	1	1	2	3	1	

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														Fı	rame	woi	rk I		-	
amino acid¹	_	2	က	4	2	9	7	8	6	10	11	12	13	14	15	16	17	18	19	20
Α					1	14			60							24	1			
В															•					
- C																				
D																				
E	1				2	1		2		64	, , , , , , , , , , , , , , , , , , ,									
F .										********				····						*********
G								58	1						64					*******
Н			2									*******		*********	} <del></del>					*******
1		2					**********	*********		••••••				•						********
K		2				********			••••••			57	64	•••••					60	••••••
L			2	59		*******	•••••		•	••••••	3	••••		******				•••••		********
М		1				*******		•••••	••••	•••••	•••••	•	•••••		••••••			*******		
· N		•		••••••		********	•			•••••		6		******						•••••
Р						••••••		*********			•••••		•	63					••••••	
Q ·	53		56		2	45				*******				-					•	
R							-		•		******	1							3	••••
S							60		3					1		40	63			
Т													*******					*******	1	•
V	2	55		1	55						61							64		64
W				••••	*********					***************************************	**********	•••••				* 1-04 Passer				
X				•••••		*******		*******						••••••		••••••		********		
Υ												********		*******				******		*******
Z	3						-					***************************************	•					*******		
-			_					_	_											•
unknown (?)																******		************		
not sequenced	11	10	10	10	10	10	10	10	6	6	6	6	6	6	6	6	6	6	6	6
sum of seq <sup>2</sup>	59	60	60	60	60	60	60	60	64	64	64	64	64	64	64	64	64	64	64	64
oomcaa³	53	55	56	59	55	45	60	58	60	64	61	57	64	63	64	40	63	64	60	64
mcaa¹	Q	٧	Q	L	٧	Q	٠\$	G	Α	Е	٧	K	K	Р	G	S	S	٧	K	٧
rel. oomcaa <sup>s</sup>	<b>%</b> 06	92%	93%	%86	92%	75%	100%	97%	94%	100%	95%	%68	100%	%86	100%	53%	98%	100%	94%	100%
pos occupied <sup>6</sup>		4			4	:	:		•	•			1	:			:	•••••••		•••••

Table 6A: Analysis of V heavy chain subgroup 1A

														CD	RI					
amino acid'	21	22	23	24	25	56	27	28	29	30	31	∢	ω	32	33	34	35	36	37	38
Α				62				1		·					41					
В			i												<u></u>					
. С		63							********											
D							1													
E									*************											
F .									69					3		3				
G				1		69	41		1		_				23					
Н										1				1			1			
								1								61	1		1	
K			63	,						1	1									
<u>L</u>															1	2				
М					•••••	********	••••••									4				
N		•••••				*********	**********			2	5						4			
P					•••••	********									1					
Q						********				••••										
<b>R</b>		1	1		*****	*******	********	•	********	1	1									7
S	63			**********	68		1			40	60			2			60			
Ţ	1			2				68		25	3				3		4			
V				·											1				69	
W									•									70		
Χ																				
Υ							27							64						
Z -																				
												70	70							
unknown (?)																				
not sequenced	6	6	6	5	2	1														
sum of seq²	64	64	64	65	68	69	70	70	70	70	70	70	70	70	70	70	70	70	70	
oomcaa <sup>3</sup>	63	63	63	62	68	69	41	68	69	40	60	70	70	64	41	61	60	70	69	
mcaa <sup>4</sup>	S	С	K	Α	S	G	G	T	F	S	S	-	-	Υ	Α	١	S	W	٧	
rel. oomcaas	%86	%86	0/08/	95%	100%	100%	29%	17%	%6	7%	<b>%9</b> 1	100%	100%	91%	59%	87%	%98	100%	%66	
pos occupied		į	ì			•		•	:	:	;	;	:			į	•	<u>:</u>	•	T

Table 6A: Analysis of V heavy chain subgroup 1A

	_			Fr	ame	wor	k II											_		
amino acid'	39	40	4		43	44	45	46	47	48	49	20	51	52	∢	В	ပ	53	54	55
Α		70									1				5					
В		<del></del>	<del></del>		<b>!</b>	<del></del>	 !		<del></del>	<del> </del>	<u> </u>	<u> </u>	<u> </u>			<u> </u>		<u> </u>	<u> </u>	
· C			<del></del>	<u></u>					<del></del>	<del> </del>	<u> </u>	<del></del>	<del></del>	<del></del>	<del> </del>	<u> </u>	<u> </u>	<del>!</del>	<del></del> -	
D		***************************************	<u> </u>	 !				1	<del></del>	<del> </del>	<del></del>	<del>:</del>	<u> </u>	ļ	ļ	<u> </u>	-	<u> </u>	<del>:</del>	
E		-	<del></del> -		<u> </u>	·······	********	69			<u> </u>	<del></del>			<del></del>	<del></del>	<del> </del>	<del></del>	<del></del>	
<b>F</b> .			•		; ;		*******	†·····			<u></u>		2				<del></del>	3	39	
G		•	1	68	(*********** !	69	*********		1	······	69	39	······		1	; !				68
Н			1	<b></b>		(	•••••	;				•••••					·····			••••••
1			<del></del>	•••••••	********	•		!					65	38			ļ	34		
K				••••			••••••							•••••••			<u></u>			
L				1		••••	68		 !	1		1						2	4	
М				••••••	••••••		••••••			67	•••••	•••••		2	********	•••••		4		
N				•••••	••••••	••••••	********	•••••	•			••••		4		•••••			22	
Р		•	68			•••••	1	••••••		••••••				••••••	44		•••••			
Q	69				69			••••	**********	••••••				********	•••••	••••••	•••••	1	1	1
R	1			1		1	******			•		4	*******	••••••		*******		1		
S					1		••••••		1	1		•••••	. <del></del>	22		•			1	1
T									*****	******			1	2	4	********		1	3	
V		·								1			2	2	16			1		
W							1		67			26		**********	***************************************	****				
X													••••••		************	**********			*******	
Υ									1							*********		20		
Z																				
•																70	70			
unknown (?)																				•
not sequenced																******				
sum of seq <sup>2</sup>	70	70	70	70	70	70	70	70	70	70	70	70	70	70	70	70	70	70	70	70
oomcaa¹	69	70	68	68	69	69	68	69	67	67	69	39	65	38	44	70	70	34	39	68
mcaa'	Q	Α	Р	G	Q	G	L	Ε	W	М	G	G	ı	ı	Р	-	-	ı	F	G
rel. oomcaa³	%66	100%	92%	97%	%66	%66	%26	%66	%96	<b>%96</b>	%66	26%	93%	54%	63%	100%	100%	49%	26%	97%
pos occupied <sup>a</sup>				3	:		:	į	i	4		4		6			1	•••••		

Table 6A: Analysis of V heavy chain subgroup 1A

	C	DR	11	·			·								·					
amino acid	26	22	58	59	9	61	62	63	64	65	99	29	89	69	20	71	72	73	74	75
А	1	34			69											43				
В																				
· C																				
D	15		1							2							70			
<u> </u>									1									33		
F				1				48				3		4						
G	1						3			67										
Н			1																	
l	4												1	44				1		
K	1		2	1			47		1		1							8		
Į.	1	1						22				2		1		3				
М														21						
N	9		59				18		••••••						••••••					
· P	1	7	•••••	•••••	•••••	•••••			•••••••		***********					*********				
Q	1	1				70			64											
R	2						2		1		69							1		
S '		1	2		1			-							5				70	
T	34	26	4		********	a 10 000 1 ba			3				66		65	24		27		67
V			••••••		•••••					1		65	3			••••••				3
W		••••		·····			•						*********	*******	***************************************					
Χ		•••••		••••				*******	•••••			••••		••••••		******		•••••		
Υ		•=	1	68						••••							•			
Z		*******																;		
•																				
unknown (?)												••••				*****				
not sequenced			<u> </u>																	
	2	70	70	70	70	70	70	70	70	70	70	70	70	70	70	70	70	70	70	70
oomcaa <sup>3</sup>	<u></u>	<del></del>	<u> </u>	<u>.</u>	<del> </del>		<u> </u>	48	*********				•••••		••••••		·			
mcaa'	····	······	÷	<u> </u>	·····	••••••	•	F			·····				*******		•••••••	••••••	S	
rel. oomcaa <sup>s</sup>	%	%	%	%	%	<b>%</b> 0	%	%69	%	%	%	%	%	%	%	%	<b>%</b> 0	47%	% 0	<b>%96</b>
	•	•		•		:			•	:	:	÷	:			:	:		:	96
pos occupied <sup>6</sup>	11	6	7	3	2	1	4	2	5	3	2	3	3	4	2	3	1	5	1	2

		•		ı	ram	iewo	rk I	11												
amino acid¹	92	77	78	79	80	81	82	۷	8	ပ	83	84	85	98	87	88	89	90	91	92
Α			64			1						3			1	70				
В																				
· C																				70
D						2							26	70						
E						64							44							
F .			·														1	1	2	
G									1				·							
Н				1				1												
		1					3	1	1								2	,		
K											3						*********			
<u>L</u>					3		63			70							2			
М					67										1		1			
N	4							1	16											
Р																				
Q				1		3														
R	3							23	1		62									
S	62		1					41	49			67			1					
T	1	69	2					3	2		4				67					
V			3				4				1						64			
W																				
Χ																				
Υ				68														69	68	
Z																				
_																				
unknown (?)																				
not sequenced																				
sum of seq <sup>2</sup>	70	70	70	70	70	70	70	70	70	70	70	70	70	70	70	70	70	70	70	70
oomcaa <sub>1</sub>	62	69	64	68	67	64	63	41	49	70	62	67	44	70	67	70	64	69	68	70
mcaa'	S	T	Α	Υ	М	Ε	L	S	S	L	R	S	Ε	D	T	Α	٧	Υ	Υ	С
rel. oomcaa <sup>s</sup>	%68	%66	91%	97%	%96	91%	%06	29%	20%	100%	99%	%96	63%	100%	%96	100%	91%	%66	% <u>/</u> 6	100%
pos occupied <sup>a</sup>		2		3	:		•	i	•		:	2	:		:	1	•	:	•	•

Table 6A: Analysis of V heavy chain subgroup 1A

		<del></del>								CDI	R III									
amino acid'	93	94	95	96	97	98	66	100	V	8	U	۵	ш	ш	9	I	_	_	×	10
Α	66	2	16		1	1	1	4	1	2	2	1	1		1	1	1	2		1
В		<u></u>	<u> </u>																	
· C					1	1	16	2		1	1	7	2	1						
D			16	5	3		3	5	4	3	4			1.	1	14	,			59
E			9				2			1			1			1				•••••
F.					1	3		2		3	1	2		2	1				28	2
G		2	14	13	20	10	14	5	20	15	16	3	3	4	15	1	1	7		
Н			į							1	1	1	-	1						
1			į	2	5	2	2		2	2	1	1			1		<u></u>			•••••
K		5	<u> </u>		2	1	.,		1								<u>.</u>			
L		1	4	4	2	5	2	1	1		4	2		1	<u></u>	<u> </u>	1		1	
М			1		2		1		1			1	1			į			10	
N				2	2	1	2	1	2	2	2	2			1	1	4			
P				20	3		1	3	2	2	2	4	2	1	4	1		1		1
Q				1			1		1	1	1									
R		55	1	5	7	8	1	4		2		1		16						
S		1	1	5	5	5	5	21	5	11	8	4	3		2	1		2		1
T	1	3	3	5	4	1	3	4	2	5	2		1			1	1			
V	3		3	2	4	3	3	3	4	2	2	2	1	2	1					
W				1	1	3	1	1			2		3				1	5	1	
Χ						•••••	•••••													••••••
Υ		1		2	3	20	5	4	9	1	2	11	20	10	6	9	10	7	1	
Z																				-
_	ļ			1	2	2	3	6	11	11	14	23	26	26	31	34	46	39	21	1
unknown (?)	ļ						,						1		1	1	******	2	3	
not sequenced			2	2	2	4	4	4	4	5	5	5	5	5	5	5	5	5	5	5
sum of seq <sup>2</sup>	70	70	68	68	68	66	66	66	66	65	65	65	65	65	65	65	65	65	65	65
oomcaa <sup>3</sup>	66	55	16	20	20	20	16	21	20	15	16	23	26	26	31	34	46	39	28	59
mcaa <sup>4</sup>	Α	R	Α	Р	G	Υ	С	5	G	-	-	-	-	-	-	-	-	-	F	D
rel. oomcaa⁵	94%	79%	24%	29%	29%	30%	24%	32%	30%	23%	25%	35%	40%	40%	48%	52%	71%	%09	43%	91%
pos occupied <sup>6</sup>	3	Ţ	•	:	:	:		•		<u> </u>	-	:	:	:		:	8	7		6

14久 SUBSTITUTE SHEET (RULE 26)

Table 6A: Analysis of V heavy chain subgroup 1A

·	Г				Fra	ame	worl	k IV		•			1
amino acid'	102	103	104	105	106	107	108	109	110	111	112	113	sum
Α		•											670
В		 !		•••••				<del></del>	<del></del>	<u> </u>	<del></del>		
С						<del></del>		<u> </u>	<del></del>	<del></del>	<del></del>		165
D		1	1				•••••	<del></del>		<u> </u>			308
E	1	1					********						297
F	2			••••				•		 !			226
G		*******	58	*******	59	1	1	•	••••••	••••••••••••••••••••••••••••••••••••••			928
Н				1			•••••		••••••	(			14
1	3		********	*******	*******	••••••	•	••••	4		••••		286
K			•	3		1		••••••	••••••		••••		325
L	.3	••••	*********	1	******		40	1			•••••		386
M	1						3			•			189
N				1									176
Р	5			•••••	••••							1	238
Q				52									494
R				1									351
S											53	51	972
TT						54	11	1	51		1		736
V	15		1				1	54		54		1	699
W		59		1									243
X													
Υ	34		1			**********							542
Z													3.
	1												578
unknown (?)													8
not sequenced	5	9	9	10	11	14	14	14	15	16	16	17	406
sum of seq <sup>7</sup>	65	61	61	60	59	56	56	56	55	54	54	53	
oomcaa <sup>3</sup>	34	*********		*******	59	54	40	54	51	54	.53	51	
mcaa <sup>4</sup>	Υ	W	G	Q	G	T	L	٧	T	٧	S	S	
rel. oomcaa <sup>s</sup>	52%	97%	95%	87%	100%	%96	71%	<b>%96</b>	93%	100%	%86	%96	
pos occupied <sup>6</sup>	9	3	4	7	1	3	5	3	2	1	2	3	

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Table 6B: Analysis of V heavy chain subgroup 1B

1					,									Fr	ame	wor	kΙ			
amino acid'	<b>,</b>	2	က	4	2	9	7	8	6	10	11	12	13	14	15	16	17	18	19	20
Α									32							34				
В																				
. C						********														
D		••••																		
E		1			5	1				35										
F																				
G								27							35					
Н			1											1						********
I									·				•	********	•••••					1
K		3	1		*****							34	33	*********	*********		*******	******	33	********
L			3	26	1										••••					••••••
M				1	1															
N																				
Р									1					33			1			••••••
Q	21		20			26														
R	1							,				1	2		•••••••					PT PT PT PT
S							27						•			1	34		•	
T									1					1		***********			2	•
V	3	21			20					************	·35			•	***************************************	********		35		34
W		•••••	•••••				•	•						**********		•				*******
X			,,,,,,,,	•••••	*********		***********				********									********
Y			•••••		•••••	******				••••••		••••	••••••			********				••••
Z				•	*******	********	•••••	•••••			•••••				•••••					•••••
-																				
unknown (?)														********			••••••			••••••
not sequenced	15	15	15	13	13	13	13	13	6	5	5	5	5	5	5	5	5	5	5	5
sum of seq²	25	25	25	27	27	27	27	27	34	35	35	35	35	35	35	35	35	35	35	35
oomcaa3									32		*********			**********	********	***********				
mcaa⁴	Q	٧	Q	L	٧	Q	S	G	Α	Ε	٧	K	K	Р	G	Α	S	٧	K	٧
rel. oomcaas	34%	34%	30%	96%	74%	%96	100%	100%	94%	%00I	%00ı	37%	34%	)4%	%00 i	%/(	%26	100%	94%	97%
pos occupied <sup>a</sup>	•		:				<u>:</u>	•	•										2	

Table 6B: Analysis of V heavy chain subgroup 1B

						•								CI	ORI					
amino acid'	21	22	23	24	25	26	27	28	29	30	31	4	ω	32	33	34	35	36	37	38
Α				30							2				6					
В																				
. C		35		·																
D											1				5		1			1
E			3						***************************************		1	******		••••	••••			.,		••••••
F		******					2		39			*****		2	2					******
G				1		40				1	14				1					1
Н									·					3	1		34			
ļ								1		1						9				
K			28																	
L									1		1					5			2	
M																23				
N							1			1	3					1	3			*****
Р		*******													.1					
Q			2								1				1		1			1
R			2					2						1						37
S	35				40			5		2	15			2	1					
T				3				32		34					1					
V				1			1			1	1				2	2			38	
W			·		,													40		
X					·					****			*******							····
Υ							36				1			32	19		1			
Z																				
-							•••••••					40	٠40							
unknown (?)																				
not sequenced	5	5	5	5																
sum of seq <sup>7</sup>	35	35	35	35	40	40	40	40	40	40	40	40	40	40	40	40	40	40	40	40
oomcaa <sup>3</sup>	35	35	28	30	40	40	36	32	39	34	15	40	40	32	19	23	34	40	38	37
mcaa <sup>4</sup>	S	С	K	Α	S	G	Υ	T	F	T	S	-	-	Υ	Υ	М	Н	W	٧	R
rel. oomcaas	100%	100%	%08	%98	100%	100%	<b>%06</b>	80%	%86	85%	38%	100%	100%	%08	48%	28%	85%	100%	95%	93%
pos occupied <sup>6</sup>	1	1	4	4		1							1		:	:				4

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Table 6B: Analysis of V heavy chain subgroup 1B

•	-			Fra	me	worl	<u> </u>													
amino acid'	33	40	41	42	43	44	45	46	47	48	49	20	51	52	4	œ	υ	53	54	55
А		39				1					1				7			1		
В		***************************************							••••••••	•			***********	************	**********	•••••				P00000000
С							***************************************			•			•••••	•••••••						
D													,	1					1	
E				1				39										1	1	
F .							2						1					1		
G				39		28					39	1			1			9	1	39
Н																		2		
I										3			34							
K					1														1	
L			1				37						1							
М										37		2	4							
N														35				20	12	1
Р		1	34				1								31					
Q	<b>3</b> 9				39		,	1												
R	1					10				į		4						3	1	
S			1			1								2				1	20	
Ţ			4											1					3	
V .														1	1					
W						********			40			33								
X			eegagaa te			*******	******													
Υ						********												2		
Z						_=									-					
		•••••	•••••													40	40			
unknown (?)						••••••												.,		
not sequenced																				
sum of seq <sup>7</sup>	40	40	40	40	40	40	40	40	40	40	40	40	40	40	40	40	40	40	40	40
oomcaa <sub>3</sub>	•••••••				·		37		***********		39	********		*******		40	40			
mcaa'	Q	Α	Р	G	Q	G	L	Ε	W	М	G	W	ı	N	Р	-	-	N	S	G
rel. oomcaas	%86	%86	85%	%86	98%	70%	93%	98%	100%	93%	%86	83%	85%	98%	78%	100%	100%	20%	20%	98%
pos occupied <sup>6</sup>	:	:	•														1		8	2

Table 6B: Analysis of V heavy chain subgroup 1B

	_	CDR		·																
amino acid'	99	22	58	59	99	61	62	63	64	65	99	29	89	69	70	71	72	73	74	75
Α	1	2			27	2				1		1				2				12
В																				
C													• • • • • • • • • • • • • • • • • • •					<del></del>		
D	1	<u>.</u>								4							35		<u> </u>	
E	2	<u>.</u>	2			1				1						1				
F.	<u>.</u>			4				39						3						
G	15		6		1					34										
Н			1	1													1			_
ı		1	1						•		********	1	1	13			•			22
· K	2	2	8				36		1	••••••				•••••		1			••••••	
L						1		1					********	1						
M													*********	23	********			1	•	1
N	17		18				1				,		*******	•	••••		4			
Р			••••••																3	
Q			•••••			36			37											
R			2				1		2		37					34		1		
S	1			2	11		1									1			37	
T		35	2		1		1						39		40	1		38		5
V .	1											38								
W											3					į				
X																				
Υ				33																
Z		·																		
_																				
unknown (?)								.,,,,,,												
not sequenced																				
sum of seq²	40	40	40	40	40	40	40	40	40	40	40	40	40	40	40	40	40	40	40	40
oomcaa,	17	35	18	33	27	36	36	39	37	34	37	38	39	23	40	34	35	38	37	22
mcaa⁴	N	T	N	Υ	Α	Q	K	F	Q	G	R	٧	T	М	T	R	D	T	S	ı
rel. oomcaas	43%	%88	45%	83%	%89	%06	<b>%06</b>	98%	93%	85%	93%	95%	%86	58%	100%	85%	%88	95%	93%	55%
pos occupied <sup>a</sup>	:	:	:	•			:	:	:	:	:	:	:	:	:		••••••	······································	••••••	

Table 6B: Analysis of V heavy chain subgroup 1B

	-			F	ram	ewo	rk II	1												
amino acid'	92	77	78	79	80	81	82	4	В	ပ	83	84	85	98	87	88	68	90	91	92
Α			35									1	2			40				
В																				
· C																				37
D	1					4							19	40			1			
E						35							19							
F			1									2							2	1
. G						1		1	2											********
H																				
1		1															1			
K											1			•••••••••••••••••••••••••••••••••••••••						
. <b>L</b>					2		39			39							2			1
М		•			37		1						-	-			2			
Ň	7							1	2											
Р												1							1	
Q																				
R	4							2	16		37									
S	27			1				35	20		1	36						1	1	
T	1	39						1			1				40					
V			4		1					1							33			
W							•													
Х																		•		
Υ				39														38	35	
Z																				
_																				
unknown (?)																				
not sequenced																	1	1	1	1
sum of seq²	40	40	40	40	40	40	40	40	40	40	40	40	40	40	40	40	39	39	39	39
oomcaa3	27	39	35	39	37	35	39	35	20	39	37	36	19	40	40	40	33	38	35	37
mcaa <sup>4</sup>	S	T	Α	Υ	М	Ε	L	S	S	L	R	S	D	D	T	Α	٧	Υ	Υ	С
rel. oomcaa <sup>5</sup>	%89	%86	98%	%86	93%	%88	%86	%88	50%	%86	93%	%06	48%	100%	100%	100%	85%	92%	%06	95%
pos occupiedº	:	:	;	:	:	:	:	:	•	:	:	:	:		1	1	:	2	:	3

Table 6B: Analysis of V heavy chain subgroup 1B

	Γ									CD	R III	<del></del>								·
amino acidi	93	94	95	96	97	86	66	100	٧	80	ပ	٥	ш	щ	ဗ	I	_	_	×	101
Α	37	1	6		1	1		2	3	1	3		1					5		
В		<del></del>	<u> </u>	<del></del> -	<del></del>	<del></del>	<del></del>			<u> </u>	<del></del>	<del></del>	<del></del>	·····	<del></del>				ļ	<del> </del>
. С		1	Ì			3		············		2	1	<del></del>	<u></u>	<u>.</u>	·	<u> </u>	<u></u>	<u> </u>	<del> </del>	
. D	<b></b>	<u> </u>	7	<u> </u>	5	2	3	1	5	4		1	·········	2	2	1	2	<del></del>	<del></del>	27
E		<del></del>	2		1	••••••		1			2		1	····	1	<del> </del>	<del> </del> -	<del></del>	<del></del>	
F			<u> </u>	1	1	3			2	1	1	1	1			<u> </u>	<u> </u>	2	15	
G		1	7	7	5	5	9	4	7	1	3	••••	2	2	1	·····	1	3		1
Н			1		••••••	*********	2		•	1	1		••••		}	••••••••••••••••••••••••••••••••••••••	<u> </u>	•	********	
1		1	<del></del>	1	1	3	1	1	1	1	1	1		•••••••			<del></del>		1	
К		1			1	********		••••••	1	1	••••••	1	*******	1	*******		1		******	•••••••
L			2	4	4	4	3		*********	1	2	1	1	2	••••••	1			2	******
M				2		1	1		***********			*******	********	*****	1				4	
N					1			1		1	1	1	••••••	********	3	•••••	1			1
P				6	4				1	1		3	2				1		••••••	
Q					1							1	2	1				·		
R	1	31		5	1	1	3					1		1				1		
S		1	3	3	1	4	3	6	3	2	2	1		1						
Т		2	1	1	2	2	1	5	1	1	1		1			1		1		***********
V	1		7	1	1		1	3	1	2		1			1	2	1			1
w			1		1		2	2		1	1	·				1		4		
X																				
Υ				5	5	4	2	3		4	3	3	2	1	2	· 5	6	2		
Z						-	į													
				1	1	4	6	8	10	11	14	20	23	25	25	25	23	18	11	6
unknown (?)																		į	3	
not sequenced	1	1	3	3	3	3	3	3	4	4	4	4	4	4	4	4	4	4	4	4
sum of seq²	39	39	37	37	37	37	37	37	36	36	36	36	36	36	36	36	36	36	36	36
oomcaa³		31			5	5	•••••••	8	10	11	14	20	23	25	25	25	23	18	15	27
· mcaa*	Α	R	D	G	D	G	G	-	-	- [	-	-	-	-	-	-	-	-	F	D
rel. oomcaa⁵	95%	79%	19%	19%	14%	14%	24%	22%	28%	31%	39%	26%	64%	%69	%69	%69	54%	20%	42%	75%
pos occupied <sup>6</sup>	•			•					12	:	14	•	•	•		•	•			5

Table 6B: Analysis of V heavy chain subgroup 1B

					Fra	ame	worl	k IV.					
amino acid	102	103	104	105	106	107-	108	109	110	111	112	113	Su
А				•									34
В											: :		
С						; :		• • • • • • • • • • • • • • • • • • •	• • • • • • • • • • • • • • • • • • •	••••••••••••••••••••••••••••••••••••••			7
D	2										**********		17
E				1							********		15
F	1							••••••	<b></b>		•••••	******	13
G			27		26		••••••		•	1	********		45
Н	1						•••••		********	41+ <del>10++</del> ++	*******	••••••	5
i l	7			•	*******	••••••		••••••	3		*******		11
K				2							•		19
L							12			1			20
M							2						14
N	1									•	*******		13
Р	1			1									12
Q				23									25
R							1						24
S	3								1		18	18	43
Ţ						21	6		16		1		39
V	6							21		18			34
W		29											15
Χ													
Υ	11	•••••				040 PBTQ Q							29
Z													
-	3												39
unknown (?)													
not sequenced	4	11	13	13	14	19	19	19	20	20	21	22	45
sum of seq <sup>2</sup>	36	29	27	27	26	21	21	21	20	20	19	18	
oomcaa³	11	29	27	23	26	21	12	21	16	18	18	18	
mcaa¹	Υ	W	G	Q	G	Τ	L	٧	T	٧	S	S	
rel. oomcaas	31%	100%	100%	85%	100%	100%	57%	100%	<b>%08</b>	<b>%</b> 06	95%	100%	
pos occupied <sup>6</sup>						:					2	1	

Table 6C: Analysis of V heavy chain subgroup 2

:				·					·		<del></del>			Fr	ame	wor	k I			
amino acid'	-	7	က	4	വ	9	7	æ	6	0	=	12	13	14	15	16	17	18	19	20
Α										3										
В																				
. С											9 6 6									
D																				
E	1					6									••••••	2				
F .							<b>-40101111</b>				,,				******					
G								6							*******	•••••				
Н															*******					
1		1													••••••					
K		*******			3								6		1					
L				6							6							6		6
M														```						
N							1													
Р							1		6					6			1			
Q	2															4				
R					2															
S							4													
T			6		1					2					5		5		6	
V		5								1		6								
W																				
X																				
Y																				
Z	3																			
-																				
unknown (?)																				
not sequenced	1	1	1	1	. 1	1	1	1	1	1	1	1	_1	1	1	1	1	1	1	_1
sum of seq <sup>2</sup>	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6
oomcaa <sup>3</sup>	3	**********	*********	6	********	6			•••••••	••••••••	6	••••••••	6	6	•••••••••			6	6	6
mcaa'	Z	٧	T	L	K	Ε	S	G	Р	Α	L	٧	K	Р	T	Q	T	L	T	L
rel. oomcaas	20%	83%	100%	100%	20%	100%	9/2	100%	100%	20%	100%	100%	100%	100%	83%	%29	83%	100%	100%	100%
pos occupied <sup>6</sup>	3	2	1	1	3	1	3	1	1	3	1	1	1	1	2	2	2	1	1	1

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Table 6C: Analysis of V heavy chain subgroup 2

•														CD	RI					
amino acid'	21	22	23	24	25	26	27	28	29	30	31	∢	ω	32	33	34	35	36	37	38
Α								1				1			1					
В																				
C		7													2		.,			
D			į									1								
E							********													
F				3			6		1											
G					.,	7					_		4		3		3			
Н																				
													1						7	
K						-														
L				2			1		6											
М														5						
N											2									
Р																				
Q										·										
R					·								2		1					7
S			1		6			6		6	2	4					4			
T	6		6	·						1	3	1								
V				2										2		7	<u> </u>			
W																<u>.</u>		7		
X								<u> </u>		<u> </u>				••••				! !	ļ	
Υ					1									•		<u>.</u>	ļ	ļ	ļ	
Z											<u> </u>	<u> </u>		-			<u> </u>			
-		<u></u>	<u>.</u>						ļ	<u>.</u>	<u></u>	<u></u>					ļ	ļ	<u> </u>	ļ
unknown (?)	<u></u>	<u> </u>	ļ	<u>.</u>				<u> </u>	ļ	<u> </u>		<u> </u>	<u> </u>			<u></u>	<u> </u>	<u> </u>	<u></u>	
not sequenced	_1	<u> </u>										<u> </u>	<u> </u>			<u> </u>	<u> </u>	<u> </u>		
sum of seq <sup>2</sup>	6	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7
oomcaa³	6	7	6	3	E	7	(	6	6	6	3	4	4	5	3	7	÷	7	7	7
mcaa <sup>4</sup>	T	C	T	F	S	G	F	S	L	S	T	S	G	М	G	· V	S	W	1	R
rel. oomcaas	100%	100%	%98	43%	%98	100%	%98	%98	%98	%98	43%	57%	57%	71%	43%	100%	57%	100%	100%	100%
pos occupied		7	2	:	:	•	1 2	:	2 2	2	•	3 4	3	:	:	1	2	1	1	1

Table 6C: Analysis of V heavy chain subgroup 2

				Fr	ame	wor	k II						Γ		-	-				
amino acid'	39	40	4	42	43	44	45	46	47	48	49	20	51	52	⋖	8	U	53	54	55
Α						6					7									
В			<u> </u>	<u> </u>					••••••		•••••			••••••			<del></del>	<u> </u>		
· C	••••••	·	<del></del>	<u> </u>	••••••		*******		••••••	••••••	•••••			•••••		<u> </u>	<del></del>	<u> </u>		
D			<u> </u>	<u> </u>		•••••	*******			*******			*****	2		<del> </del>	<del> </del>		3	6
E			<u> </u>				•	7	7					•••••			<del></del>			
. F .						•••••••	**********		•••		•••••	••••		2	••••••					
G		1		7		1		••••	*******			*******		*********	••••••					
Н							*********					2	********	******	********					1
1		***********				•			•••••				6							
κ		•••••			6		•••••			***************************************					••••••	•••••	•••••			
L		*****	• • • • • • • • • • • • • • • • • • •				7			7		2	1	1	*******	••••	•••••	·		
M									**********	***************************************			••••••	*******	•••••	******				
N									***********		•				********	•••••	•••••		3	••••••
Р		5	7						**********	•••••••				••••••••	*****	•	•	•		
Q	6							•••••	••••••••	********	•••••••••••••••••••••••••••••••••••••••	•••••••		••••••	***************************************		••••			******
R	1				1				,			2			••••••					
S		1									į						******	2		
T							·													
V										·	į									
W									7			1						4		
X														1				1	1	
Y														1	1					
Z -																				
_															6	7	7			
unknown (?)																				
not sequenced																				
sum of seq²	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7
oomcaa³	6	***********	7	7	6	6	7	7	7	7	7	2	6	2	6	7	7	4	3	6
mcaa•	Q	Р	Р	G	K	Α	L	Ε	W	L	Α	Н	ı	D	-	-	-	W	D	D
rel. oomcaa <sup>s</sup>	96%	71%	100%	100%	%98	%98	100%	100%	100%	100%	100%	29%	%98	29%	%98	100%	100%	57%	43%	%98
pos occúpied <sup>6</sup>	2	3	1	1			1	1	1	1	1		2	•		:	:	•	:	2

Table 6C: Analysis of V heavy chain subgroup 2

		DR	11																_	_
amino acid'	26	22	58	29	09	61	62	63	64	65	99	<b>6</b> 9	89	69	70	71	72	73	74	75
Α																				
В						*******	******		•	••••••					•••••			•		
. С			••••••		******	••••••	*****					••••			••••	•••••••				
D	5		••••••				******	••••		***********	•		•••••••		••••		6	1		
E	1			•••••	********	10000000	********	••••••	1			•	•••••		********		••••			
F		1	••••••	1	•	•••••	•••••	•••••		***********			••••••••		••••					
G			•	**********	•••••	********				••••••			•••••••••••••••••••••••••••••••••••••••	•••••••••						
Н				1	••••	****				••••••	********	•••••••••••••••••••••••••••••••••••••••	•••••••	•••••••••••••••••••••••••••••••••••••••	••••••				*******	
l						*******			***************************************		•••••••••••••••••••••••••••••••••••••••	•		6	**********			•		
К	1	6							4			•••••••••••••••••••••••••••••••••••••••			•••••••••••••••••••••••••••••••••••••••	6	**********	**********		6
L								7				7								
М.																		*******		
N						•									***************************************		1	••••••	:	
. Р						2														
Q																				
R			2			1			2		7					1			•	1
· S			2		6		7			4		·	1		5				7	
Т						4				3			6		2	, ,		6		
V				*******										1						
W		,		1			,													
X					1	*******														
Υ			3	4															į	
Z																				
			••••••			*******	******													
unknown (?)						••••••														
not sequenced																				
sum of seq²	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7	7
oomcaa3	5			4		4	•••••	7	4		7	7	6	6	***************************************		6	6	7	6
mcaa <sup>4</sup>	D	K	Υ	Υ	S	T	S	L	K	S	R	L	Ţ		S	K	D	T	S	K
rel. oomcaas	71%	%98	43%	57%	%98	57%	100%	100%	57%	57%	100%	100%	%98	%98	71%	%98	%98	%98	100%	%98
pos occupied <sup>6</sup>	3	2	3			:	:	: :	:	:	1	1	2	2	2	2				

Table 6C: Analysis of V heavy chain subgroup 2

•					Fran	1614	nrk I	111												
amino acid'	9	7	<b>&amp;</b>						~		83	4	-2	9	_	<u> </u>	6	_		7
	_	_	<u> </u>	7	<u> </u>	<u> </u>	<u> </u>	_		<del>-</del>	<u> </u>	· α	<u></u>	<u> </u>	<u> </u>	88	<u></u> &	<u>ர</u>	:	92
Α	ļ	<u> </u>	<u> </u>	<u> </u>	<u></u>				<u>.</u>	<u> </u>	<u> </u>	<u></u>	1		<u> </u>	5	<u></u>	<u> </u>	<u> </u>	
B	<b>.</b>	<u> </u>	ļ	<u> </u>	<u>.</u>		<u>.</u>	ļ	<u> </u>	<u>:</u>	<u> </u>	<u></u>	<b></b>		ļ	<u> </u>	<u></u>	<u> </u>		
. C		<u> </u>	<u> </u>	<u> </u>	<u>.</u>	<u></u>	<u> </u>	<u></u>	<u> </u>	ļ	ļ		<u></u>			<u> </u>		<u> </u>		7
<u>D</u>	-	<u> </u>	<u> </u>	<u> </u>			ļ				6			7		ļ		ļ		
E	<u> </u>	<u> </u>	ļ	<u>.</u>	<u> </u>		<u> </u>	<u> </u>	ļ			••••••				<u> </u>		ļ		
F .	ļ	ļ. 	<u> </u>	ļ	1		······		ļ								*****			
G	ļ	<u> </u>	ļ					<u></u>	<u></u>					•••••		2				·
<u> </u> H	<u></u>	<u></u>	ļ		••••••			<u>.</u>	<u>.</u>											
	<u></u>	<u>.</u>				2		1	<u>.</u>											
K	<b></b>	<u> </u>			•															
L L					6															
M							7			5								·		
N	5								6		1									
Р												7								
Q		7																		
R				-											*********		***************************************			
S	2		·									•						•		
Т						5	********	5						•••••••••••••••••••••••••••••••••••••••	7	·····	7		i	
V			7	7	•••••••••••••••••••••••••••••••••••••••		•••••	•••••		1	Ī	•••••	6	*****		<u>†</u>			i	*****
W					********		•••••				······				••••••	<del></del> †			·····-	
Х					***************************************	•••••••			•••••					••••••						
Y					•••••••••••••••••••••••••••••••••••••••	•••••••		••••	•••••							<del></del>		7	7	
Z					•	•••••••	•••••		**********							-				
_								1	1	1						i		i	T	
unknown (?)					***************************************	********	**********		*******			••••••	******	•••••						-
not sequenced			Ī			*********		••••••		<u>-</u>			*****			·i			<u>-</u>	
sum of seq <sup>2</sup>	7	7	7	7	7	7	7	7	7	7	7	. 7	7	7	7	7	7	7	7	7
oomcaa³	5	7	7	7	6	5	7	5	•••••	<del>:</del>	6	7	6	7	7	·····i	7	7	7	7
mcaa'	N		•••••••••••••••••••••••••••••••••••••••	٧	L	T	М	••••••	N			Р	٧	D	T	Α	T	Y	Y	С
		%	્ર	્ર			%					્ર		e S	و		 Q	Q	o Q	 Q
rel. oomcaas	71%	100 1	100%	100%	%98	71%	001	71%	%98	71%	%98	100%	%98	100%	100%	71%	0001	100%	<b>%00</b>	%00 <b>1</b>
pos occupied <sup>6</sup>	······	······i	·····	·····	•	••••••	•••••	······ <u> </u>	•	:	2	:	•	•••••••••••••••••••••••••••••••••••••••		:	1	•••••		
• ;	········	ā	i.	i.					<u>-</u> :	ž	<u>-</u> .;.		<u>-</u> i.	!		i.	!	!		!

Table 6C: Analysis of V heavy chain subgroup 2

									<del></del>	CDI	R III									
amino acid¹	93	94	95	96	97	86	66	100	<	В	ပ	۵	ш	ட	g	Ξ	_	_	×	101
Α	5							1	2	1										
В																				
С																				
D																				6
E								2			1									
F						*******	**********			•••••		•••••					**********		3	1
G						1	1		1	2	1	1	1	1	********		4			
Н		1		1						•••••			••••••••		*******		4			
l			3			2							•	***************************************						
К			•••		***************************************		1	••••••					••••••							
L						•	•••••	1		1	•••••••		*******	********					1	
M.			**********			••••••	•	1							*********		•••••		2	
N		********		1	2		•										1			**********
Р				1	1	••••••	1		1											
Q		********	1	·			•••••		************						•••••••					
R		6	1			1	*******	•••••••••••••••••••••••••••••••••••••••	1	*******						••••••				
S		**********		1	*******	1	1		***************************************	•••••••••••••••••••••••••••••••••••••••					•••••	•••••••				
Т				1	*******	••••••	1		1					•		•••••••••••••••••••••••••••••••••••••••			Ī	
V	2		1	1	1	••••••	1	1	********		. 1				********		İ			
W			•••••			1	•	-							1			1		**********
X				•••••••	***************************************				*********									*******		*******
Y				*******	2	••••••				••••	1	2	1	1	1		•	2		
Z					*********		*****						•••••••••••••••••••••••••••••••••••••••					*********		
<del>-</del>										2	2	3	4	4	4	6	5	3		
unknown (?)																				
not sequenced			1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1
sum of seq²	7	7	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6
oomcaa³	5	6	3	1	2	2	1	2	2	2	2	3	4	4	4	6	5	3	3	6
mcaa*	Α	R	1	Н	N	I	G	E	Α	-	-	-	-	-	-	_	-	-	F	D
rel. oomcaas	71%	%98	20%	17%	33%	33%	17%	33%	33%	33%	33%	20%	67%	9/2/9	9/2/9	100%	83%	20%	20%	100%
pos occupied <sup>a</sup>							:					: :								1

Table 6C: Analysis of V heavy chain subgroup 2

		· ·			Fr	ame	wor	k IV				·	]
amino acid'	102	103	104	105	106	107	108	109	110	111	112	113	sum
А									1				35
В						·····					<u> </u>		
С						·	<u> </u>	<u></u>			<u> </u>		16
D				: :			<u> </u>	<u> </u>	<u> </u>	<u> </u>			43
E			-		·			<u> </u>	<u>.</u>	ļ		•••••	21
F						<del></del>	······································	<u> </u>	••••••••••••••••••••••••••••••••••••••		<u> </u>		18
G			6		6	}	•	<del></del>	ļ	<del></del>			55
Н					;*************************************	}	)		<u> </u>				6
					•	-		·		<del></del>			29
K		<u>.</u>		1			1			<del></del>		•••••	42
L	1			,			3					******	78
M		: :				••••••	•••••	•••••		••••••			20
N				•••••	•	***********							23
Р	1						1		*******	•••••			41
Q				3									23
R				2									41
S											6	3	82
T						6	1		5		·		102
V	3							6		6			68
<u>W</u>		6		*******									29
X				********	*********								4
Υ	1					•••••							35
Z													3
				•••••									56
unknown (?)													
not sequenced	1	1	1	1	1	1	1	1	1	1	1	4	54
sum of seq <sup>7</sup>	6	6	6	6	6	6	6	6	6	6	6	3	
oomcaa <sup>1</sup>	3		•••••••••••••••••••••••••••••••••••••••	•••••••••••••••••••••••••••••••••••••••	•••••••	•••••••••	3	6	5	6	•••••••••••••••••••••••••••••••••••••••	3	
mcaa⁴	٧	W	G	Q	G	T	L	٧	T	٧	S	S	
rel. oomcaa <sup>s</sup>	20%	100%	100%	20%	100%	100%	20%	100%	83%	100%	100%	100%	
pos occupied <sup>6</sup>	4	1	1	3		1 62	4	1	2	1	1	1	

WO 97/08320 .

PCT/EP96/03647

Table 6D: Analysis of V heavy chain subgroup 3

į								-	<del></del>					F	rame
amino acid'	-	7	က	4	2	9	. 7	8	6	10	=	12	13	14	15
Α					1		1			12		1		3	1
В	-		1			1							1		
· C															
D	1					1				16	,				
E	110		9		15	166			9				8		2
F .											4				
G								181	193	174		1			202
. Н			5										4		
1												9			
K		5	3									·	26		
L		1	5	176	43						140			1	
М		12		1										·	
N										1					
Р													1	194	
Q	41		138	1	3	12							162		
R			6										4		
S							178			2				8	
T							1								
· V	5	147		1	118						62	195			
W															1
X													.,		*******
Υ								**********		***********					
Z	8										-				
											********	************			
unknown (?)				********				444444	***********	*********					***********
not sequenced	47	47	45	33	32	32	32	31	10	7	6	6	6	6	6
sum of seq <sup>2</sup>	165	165	167	179	180	180	180	181	202	205	206	206	206	206	206
oomcaa <sup>3</sup>	110	147	138	176	118	166	178	181	193	174	140	195	162	194	202
mcaa⁴	Ε	٧	Q	L	٧	Ε	S	G	G	G	L	٧	Q	Р	G
rel. oomcaas	67%	9068	83%	%86	%99	92%	<b>%66</b>	100%	<b>%96</b>	85%	%89	95%	79%	94%	%86
pos occupied <sup>6</sup>	5			:			:	:			:	:	•	:	•

Table 6D: Analysis of V heavy chain subgroup 3

,	work	I					·								
amino acid'	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30
Ä								183	192		1				
В															
· C						1	209								
D															7
Ε	8			********				8			3		1		
F		1	1		***********	1	• • • • • •	•••••				201		201	·
G	134	******	**********				• • • • •		2		207				3
Н		***********	*************					**********						-	1
		**********						2				3	17	1	
Κ				15						·					4
L			205		201		,					6		3	
M			1										1		
N													10		10
Р		***********				•		1					2		
Q		**********	1		·				,						
R	62			191	*********										11
S		206			•••••••	207		4	2	209			15		174
T	4	1		2	•••••••	****		4	4			1	163		
V					8	********		7	9				1	6	
W			***************************************			••••••	***************************************	******		•••••					
X									*******	••••					
Υ							•••••••			***************************************	***************************************	••••••••••			
Z															
					•••••	•••••					•	************			
unknown (?)															
not sequenced	4	4	4	4	3	3	3	3	3	3	1	1	2	1	2
sum of seq²	208	208	208	208	209	209	209	209	209	209	211	211	210	211	210
oomcaa³	134	206	205	191	201	207	209	183	192	209	207	201	163	201	174
mcaa⁴	G	S	L	R	L	S	С	Α	Α	S <sub>.</sub>	G	F	T	F.	S
rel. oomcaas	64%	%66	<b>%66</b>	92%	<b>%96</b>	%66	100%	%88	92%	100%	%86	95%	78%	95%	83%
pos occupied <sup>6</sup>	4	•••••••••••••••••••••••••••••••••••••••					•••••••				3				

Table 6D: Analysis of V heavy chain subgroup 3

				CE	RI									F	rame
amino acid'	31	⋖	8	32	33	34	35	36	37.	38	39	40	41	42	43
Α	1			17	80		1			1		187		1	
В				·											
· C												1		1	
D	26			3	7		2								
E	1				10									1	1
F.				5			,	·							
G	13				31		1					2		209	-
Н		***********		4			88								
	1			1		15			12						
K	7										1				202
L	3					3			2	3	1	2	1	•	******
M						193		-			٠		*********	••••••	
N	35			8	3		34	***********	***********	•	*******				*********
Р				1			1	***************************************	*************	•	*********	4	191	••••••	2 200 200 ann a
Q				*************************	••••••••••••	••••••		***************************************	**********	**********	209		1		1
R	7							***********		207		7	*******		8
S	103			17	8		72				•	3	14		*********
T	9				15		10					4	5		***********
V	2		e.		7	1			197			2			******
W					30	,		212							***********
Х	1													·	
Υ	1			154	19		3						•		**********
Z															
-		210	210												
unknown (?)															
not sequenced	2			2	2				1	1	1				
sum of seq <sup>2</sup>	210	210	210	210	210	212	212	212	211	211	211	212	212	212	212
oomcaa,	103	210	210	154	80	193	88	212	197	207	209	187	191	209	202
mcaa*	S	-	-	Υ	Α	М	Н	W	٧	R	Q	Α	Р	G	K
rel. oomcaas	49%	100%	100%	73%	38%	91%	42%	100%	93%	%86	99%	988%	%06	%66	95%
pos occupied <sup>6</sup>	•••••	1	1	9	•••••••••••••••••••••••••••••••••••••••										

Table 6D: Analysis of V heavy chain subgroup 3

	work	11							_						
amino acid	44	45	46	47	48	49	20	21	52	<	æ	U	53	54	55
Α	1					77	42		1	2		14		7	
В			3			·				1					
· C	·												1		
D			1							7			94	8	3
E			198	• • • • •					3	2	1		2		1
F		· · · · · · · · · · · · · · · · · · ·	- - - - - - - - - - - - - - - - - - -			• • •	7	1	2	1				1	8
G	207		• • • • • • • • • • • • • • • • • • •	· · · · · · · · · · · · · · · · · · ·		33	11		10	46			4	163	85
Н					<u>.</u>		6			1					
					3	·	3	191		1					1
K								1	37	2	30		3	1	
L		211			5		12	1							
М		•••••					1	1					,		
N		•••••		••••••			13		7	9	2		13	11	1
Р		1	*******	*********						1			1		
Q		************	7	**********	••••••		7			10					
R	1	***********		******			24	1	17	5	1		2		16
S	3		********	1		102	11	9	118	43		1	74	17	82
T	) · · · · · · · · · · · · · · · · · · ·			**********			3	5	4	2		13	12	3	3
- V			3		204	, 	49	2	*********	1		6			
W			****	210	1000-100000	0 1000000 1000	1		8	6		·			
X			********	********	***********								4		3
Υ				1	**********	*******	22		5	58					8
Z				-											
-					***********		*********	************	**********	14	178	178	2	1	1
unknown (?)					••••••	41 <b>44</b> 4 <b>44 44</b>					**********				
not sequenced															
sum of seq <sup>2</sup>	212	212	212	212	212	212	212	212	212	212	212	212	212	212	212
oomcaa³	207	211	198	210	204	102	49	191	118	58	178	178	94	163	85
mcaa'	G	L	Е	W	٠٧	S	٧		S	Υ	_	-	D	G	G
rel. oomcaa⁵	0/86	100%	93%	%66	<b>%96</b>	48%	23%	%06	26%	27%	84%	84%	44%	77%	40%
pos occupied <sup>6</sup>	4	2					:			:		:	:		•••••••••••••••••••••••••••••••••••••••

Table 6D: Analysis of V heavy chain subgroup 3

	(	CDR I													
amino acid'	26	57	58	59	09	61	62	63	64	65	99	67	89	69	20
Α	9	1	2		174	33							1		
В	1	2													
· C															
D	11		17			160									***
E	8	3	2			1			2						
F .	1		3	2								207			
G .	5	1	5	•	4	5				212	1				
Н	1		4												••••••
l	3	37	2					8					14	208	
К	1	61							199		8				
L	1	1	1		1							1		1	
M	8		2		1										
N	51		4			2			. 2						
Р	1	1			6	8	18		1						••••••
Q	3	2							2		2	*************			••••••••
R	5	4			5	••••••		***************************************	6	•••••••	201			••••••	••••••
S	48		11		4		193	•••••••••••••••••••••••••••••••••••••••			•	2	7		211
Т	42	97	5		7								189		1
, V		2			10	2		204				1		3	•••••
· W			2		•				•••••	*************	***************************************	*****	***************************************		
Χ	4		1	***************************************		1	**********	***************************************	•••••••••••••••••••••••••••••••••••••••	•••••••••••		***************************************	,		*******
Υ	9	•	151	210	***************************************	••••••••	1	***********	***************************************	***************************************	•••••••••••••••••••••••••••••••••••••••	1	1		
Z	***********			•••••		************		************		••••				••••••	**********
_															
unknown (?)															
not sequenced											,				
sum of seq²	212	212	212	212	212	212	212	212	212	212	212	212	212	212	212
oomcaa,	51	97	151	210	174	160	193	204	199	212	201	207	189	208	211
mcaa⁴	N	T	Υ	Y	Α	D	S	٧	K	G	R	F	T	i	S
rel. oomcaas	24%	46%	71%	<b>%66</b>	82%	75%	91%	%96	94%	100%	95%	98%	%68	%86	100%
pos occupied <sup>6</sup>	:	12					•			•••••					

Table 6D: Analysis of V heavy chain subgroup 3

					· · · · · · · · · · · · · · · · · · ·					Frar	newo	rk III	·		
amino acid'	71	72	73	74	75	92	77	78	79	80	81	82	4	8	ပ
Α				57			1	8						1	
В											2				
· C									**************************************		**************************************				
D		199	38		2	2			1			•	10		
E		6	,		4		**********	*********			5	***************************************		**************************************	
F								•	13						
G													.1	4	
Н						1			1		2		2		
l			1				2	2	,			3	1	1	
K					186	6							3		
L								188		209		3	1		212
M	1				2		10	3			~	205	•		
N		5	170		2	188					3		181	10	
Р							1								
Q					7						199				
R	211			,	1	1							2	8	
S				153	8	10	56		3				6	186	
Ţ							142				1		4	2	
V				1				11		1		1			
W															
Χ		2	2			4							1	·	
Y			•••••				****		194						
Z						·									
-							******			******	******************************				
unknown (?)						***********									
not sequenced			1	1	·										
sum of seq'	212	212	211	211	212	212	212	212	212	212	212	212	212	212	212
oomcaa³	211	199	170	153	186	188	142	188	194	209	199	205	181	186	212
mcaa¹	R	D	N	S	K	N	Ţ	L	Υ	L	Q	М	N	S	L
rel. oomcaa <sup>s</sup>	100%	94%	81%	73%	%88	9%68	67%	%68	92%	<b>%66</b>	94%	92%	85%	98%	100%
pos occupied <sup>a</sup>	2			3					•						•••••

Table 6D: Analysis of V heavy chain subgroup 3

•															<del></del>
amino acid'	83	84	82	98	87	88	68	06	91	92	93	94	95	96	97
Α		149	1		1	207					173	2	15	9	11
В															
· C									1	210		5	2		1
D		5	15	209								2	54	7	6
· E	1		190										11	2	11
F							1		15			1	·	9	6
G	1	1	6			4	1				2	8	34	26	35
Н		1							1					3	11
		8					2						4	15	10
К	30											60	- 4	3	5
L							18					1	6	11	7
М					2		1							6	1
N		1		1								2	20	4	3
Р		9									1	3	4	29	10
Q				1								5	3	9	2
R	177											103	9	30	19
S		1			1							3	9	8	11
T	3	28			207		1				25	15	7	6	20
V		9					187				10	1	. 7	7	15
W			**********							1			3	4	3
X			**********	1	***********			********							
Y							*********	211	194				12	9	8
Z .															
-		4					*********	••••••	**********	*********			1	3	4
unknown (?)					***********	*******	**********		*********	••••	*********	1010-			•••••
not sequenced					1	1	1	1	_ 1	1	1	1	7	12	13
sum of seq <sup>2</sup>	212	212	212	212	211	211	211	211	211	211	211	211	205	200	199
oomcaa <sub>3</sub>	177	149	190	209	207	207	187	211	194	210	173	103	54	30	35
mcaa'	R	Α	E	D	T	Α	٧	Y	Υ	С	Α	R	D	R	G
rel. oomcaas	83%	. %02	<b>%</b> 06	%66	%86	%86	%68	100%	92%	100%	82%	49%	26%	15%	18%
pos occupied <sup>a</sup>	5	10	4	4	4	2	7	1	4	2	5	14	18	20	21

Table 6D: Analysis of V heavy chain subgroup 3

					CD	R III									
amino acid'	86	66	100	⋖	മ	ပ	Q	m	u.	9	Ξ	_		×	101
Α	7	13	7	9	6	2	3	5	5		9	:	13		2
В															
· C	13	5		1	2	11	3		2					1	
D	11	7	10	4	2	3	10	3	3	1		3	2		146
E	6	3	1	13		1	1								1
F	3	5	4	5	5	6	3	5	7	2		1	1	65	1
G	34	17	35	17	14	23	10	5	1	5	3	2	32		6
Н	3	4	3	2	9	2		1	3	1	2	8	1		
l	6	11	4	4	3	1	3	10	3	3	2		1	2	
K	2	11			3	1									
L	26	13	4	12	8	· 2	6	3	10	3				2	1
M		1	2								1			32	
N	4	6	4	3	2	2	6				2	5			2
Р	6	5	5	6	9	8	2	3	2	1		3		9	
Q	4		1	1	1	. 1	. 1					1			
R	4	10	9	7	5	5	2	3	1		1		2		4
S	16	28	27	25	24	8	11	9	3		2	3	1	1	1
T	6	12	9	17	17	1	2	5	1	9	3	1			
<u> </u>	13	7	15	4	3	6	2	12		1	1	1	1		
W	6	5	6	7	2	4	***********			1	***************************************	6	10	******	
X		***********		1			************								1
Y	16	14	17	5	8	18	20	13	20	25	28	32	28		
Z _															
_	12	21	35	54	73	87	102	110	126	135	134	120	91	71	21
unknown (?)							3	2	1	1			3	2	
not sequenced	14	14	14	14	15	19	21	22	23	23	23	25	25	. 26	25
sum of seq?	198	198	198	197	196	192	190	189	188	188	188	186	186	185	186
oomcaa <sub>1</sub>	34	•••••	35	54	73	87	102	110	126	135	134	120	91	71	146
mcaa•	G	S	G	-	-	-	-	<u>-</u>	-	-	-	-	-	-	D
rel. oomcaas	17%	14%	18%	27%	37%	45%	54%	58%	67%	72%	71%	65%	49%	38%	78%
pos occupied <sup>6</sup>	20	20	19	20	:	•••••••••	•••••••••••••••••••••••••••••••••••••••	•••••••••••••••••••••••••••••••••••••••			•••••••••••••		:		•

Table 6D: Analysis of V heavy chain subgroup 3

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					Fr	amew	ork l	V					ı
amino acid'	102	103	104	105	106	107	108	109	110	111	112	113	su
Α	1		1			2		ĺ					17
В			·····	1			<del>-</del>						
С	•							•••••••					4
D	2	······································	Ī				į						11
Ε					1								8
F	2												8
G			140		130		1						27
Н	4												1
	15			<u></u>	<u></u>				1	1			6
K		<u></u>	<u></u>	13			<u></u>						9
L .	10	<u></u>	<u></u>	1			91					2	18
. М	į	<u></u>		<u></u>	<u></u>		6						4
N	1					1							8
Р	17					1	1						5
Q		<u> </u>		111									9
R		<u> </u>		8			<u> </u>						14
S	7	1								<u></u>	118	110	30
T						123	27		122			1	14
<u> </u>	34		1			1		125		119		.,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	18
W		158											6
Χ													
Y	82												15
Z													
	9	2	2	2	2	2	2	2	2	2	1	1	20
unknown (?)				**********									
not sequenced	27	50	67	75	78	81	83	84	86	89	92	97	16
sum of seq <sup>2</sup>	184	161	144	136	133	130	128	127	125	122	119	114	
oomcaa <sup>3</sup>			**************	111				125		**********			
mcaa4	Υ	W	G	Q	G	T	L	V	T	V	S	S	
rel. oomcaas	45%	%86	97%	82%	%86	95%	71%	%86	98%	%86	%66	<b>%96</b>	
pos occupied	12	3	:	6	3	:		2	3	3	:	:	

Table 6E: Analysis of V heavy chain subgroup 4

														F	ram	ewo	rk I			
amino acid'		7	က	4	5	9	7	8	့ တ	10	11	12	13	14	15	16	17	18	19	20
Α									19					1			1		1	
В															•		<u> </u>	<u>.</u>		<del></del>
. С								·······	<u> </u>		<u> </u>					<del></del>	<del></del>		<del></del>	<del></del>
D								<u> </u>	<del></del>	•••••	Ì					<del></del>	<del> </del>	 !	<u></u>	<del></del>
E				•		32			<u> </u>	•••••••						44			<del></del>	
F .										••••		•					·	•••••		•
G			·				}	54	1	53		_	······································			2		********	••••••	•
Н		•	4	•	2		}	••••••••••••••••••••••••••••••••••••••		•••••					}			•••••	••••	•••••
			•	······	 !			; ;	•	********		*******		••••••					••••••	
К			••••••									1	54	••••••			•••••	*******	1	
L		7		54		•••••	••••••				53	19	••••••	1				53	•••••	50
M			•••••				•••••						•••••							
N		•••••	•••••		•••••	•••••	••••			********				********	••••••	•••••			•••••	
Р		•••••		•	********	••••••		••••••	33		•	••••••	•••••••	51	1	*********				2
Q	52	•••••	50	*********	51	20	•			*****			••••••	*******	•••••	7				
R	1				********	•••••			••••••			**********		********	*******					
S.		·				•	33						*********	••••••	52				52	******
T						******		*******	1		İ						52	*********		
V		47				1			<u> </u>			34	*******		*********			*******		1
W				•••••	*****	*******	20		·····						*********	•••••••••••••••••••••••••••••••••••••••				••••
Х				******					i	·····	•		******	***********	•••••					•••••
Υ		**********				********			•		*****			•••••	***************************************					••••••
Z	1								<del>-</del>		******		*******			-				
_												i								
unknown (?)						••••••	********	***************************************					•••••••••••••••••••••••••••••••••••••••	•••••••••••••••••••••••••••••••••••••••			••••••			•••••
not sequenced	3	3	3	3	4	4	4	3	3	4	4	3	3	4	4	4	4	4	3	4
sum of seq <sup>2</sup>	54	54	54	54	53	53	53	54	54	53	53	54	54	53	53	53	53	53	<del></del>	
oomcaa³	• •	•	÷		:		:		33	:		•	•						**********	*******
mcaa⁴	• •	٧	•		Q				······÷	G	L	٧	K	Р	S	Ε	T	L	S	L
rel. oomcaa⁵	%96	87%	93%	100%	%96	%09	62%	100%	61%	100%	100%	63%	100%	%96	%86	83%	%86	100%	%96	94%
pos occupied <sup>a</sup>	3	•		1				<del>-</del>	•••••	1	<u>-</u>	3	:	3	:	•	2	1	:	3

Table 6E: Analysis of V heavy chain subgroup 4

														CD	RI					
amino acid'	21	22	23	24	25	56	27	28	29	30	31	⋖	<b>&amp;</b>	32	33	34	35	36	37	38
Α			22											1						
В																				
· C		53													1					
D		*******	1	********							4	1	1	1			1			
<u> </u>																				
F					1				22					1	1				1	
G						53	53				21	3	4				8			
Н							1							2						
<u> </u>			1					1	32										51	
K															*******					
<u> </u>																			1	••••
M																				•
N										1	1		2	2			1			
P								3												
<u> </u>											1									
R		••••••				1				3	2		1							57
<u>S</u>			2		35			51	1	52	25	5	9	1			44		1	
T	53	*******	29								2	1					3			••••••
V		••••••		55		1			1										3	
W							•					1			2	56		57		
X				••••••			•••••	•••••												•••••
Υ					19		1							48	52					
- Z																				
-					*******		400000					45	39							
unknown (?)	ļ					••••••									*******					
not sequenced	4	4	2	_ 2	2	2	2	2	1	1	1			1	1	1			-	
sum of seq <sup>2</sup>	53	53	55	55	55	55	55	55	56	56	56	56	56	56	56	56	57	57	57	57
oomcaa3	•	÷	29	******	**********		·····		32	*********		45	39						51	57
mcaa'	T	С	T	٧	S	G	G	S	1	S	S	-	-	Υ	Υ	W	S	W	1	R
rel. oomcaas	100%	100%	53%	100%	64%	<b>%96</b>	<b>%96</b>	93%	57%	93%	45%	80%	20%	<b>%98</b>	93%	100%	77%	100%	89%	100%
pos occupied <sup>6</sup>	•	•	5	:	•	i	•	•	•						•		5	1		-

Table 6E: Analysis of V heavy chain subgroup 4

										-			,							
				Fr	ame	wor	k II													
amino acid <sup>1</sup>	39	40	41	42	43	44	45	46	47	48	49	20	51	52	⋖	8	ں	53	54	55
Α			8	1							1									
В																				
· C																		<u> </u>		
D													<del></del>	1				1		
Е				1				56				22		···········				 !		
F .												1		1	<del></del>	·	•	 !		
G				55		55		}			56	1		*********		<del></del>	•	1		57
Н		2							9			******				••••••		24		•••••
I									**************************************	54		1	54		********					•••••••
К				*********	54		••••••		·								••••••	*******		••••••
L		1		••••••	*********		55			2					••••••	•••••	*******			*******
. М								•	********			********	*******	*******	••••••	•••••				
N			*********			•••••	•••••	••••				•••••		21	•••••••		*******	•••••		••••
Р		50	49		******		2		********				••••••	•••••		••••		•••••		*******
Q	56	********		**********	******		*********	1	********			1	*******	********	•••••		******	********		
R	,			•	3	2	******	********				9		1				•••••		•••••
S		3		••••••	•			*******			·	7	*******	1	*******				52	********
Т	1	1						••••							*******			8	•••••••••••••••••••••••••••••••••••••••	••••••
V									••••••	1			3			**********				
W					•••••			en een ue ee	56						*******		••••••			•••••
Х							***************************************		••••			*******								********
Υ									1		i	15	******	32	***************************************	••••••		23		******
Z												***************************************	· <del></del>		***************************************		********			******
-						٠					i				57	57	57			
unknown (?)					••••••															*******
not sequenced								******									•••••••••••••••••••••••••••••••••••••••			•••••
sum of seq²	57	57	57	57	57	57	57	57	57	57	57	57	57	57	57	57	57	57	57	57
oomcaa¹			:	:	:	:	:			:	56					***********			*******	*******
mcaa•	Q		Р	G	!	G	L				G		1	Υ	-	-	-	**********	S	••••••
rel. oomcaas	%86	88%	%98	%96	95%	%9€	%9€	%86	%8(	95%	%86	%61	92%	26%	100%	100%	100%	42%	91%	100%
pos occupied <sup>a</sup>			:			:	:			•	<u>გ</u>		თ 2				:	:	ი 2	

Table 6E: Analysis of V heavy chain subgroup 4

	C	DR											ı							
amino acid'	26	22	58	59	09	61	62	63	64	65	99	29	89	69	70	71	72	73	74	.75
Α		1									1		1			1				1
В			******																	
· C		•																		******
D			2									1		*********	•••••		55			
<u>E</u>										*******				*********			1			
F		•••••		3														1		
G	1		•••••							1										
Н			2																	••••
	1	1										1	1	48		3				•••••
K					1				53									1		51
<u> </u>						1		55				1				3				1
M														7				2		
N	2		40		53								2							1
Р						54		1												
Q		••••••															1			
R	2								3		56									2
S	49		1		2		56			56			1		56			1	57	
Ţ	1	54	1			1			1				51		1			52		
V	1	1										53		2		50				1
W																				
X										*******	*****			******				******		
Υ			11	54																
Z													~							
_					****	••••••								******	•••••					
unknown (?)						*******												•••••		
not sequenced					1	1	1	1				1	1							
sum of seq <sup>2</sup>	57	57	57	57	56	56	56	56	57	57	57	56	56	57	57	57	57	57	57	57
oomcaa3	·	····	÷	*********	******	********	**********	••••••	••••••	• • • • • • • • • • • • • • • • • • • •	56		**********	*******	56	50	55	52	57	51
mcaa*	S	T	N	Υ	N	Р	S	L	K	S	R	٧	T		S	٧	D	T	S	K
rel. oomcaa <sup>s</sup>	96%	95%	70%	95%	95%	<b>%96</b>	100%	%86	93%	%86	%86	95%	91%	84%	%86	%88	<b>%96</b>	91%	100%	%68
pos occupied <sup>6</sup>	:	:	:			:	:	:	•	:	:	:						5		

Table 6E: Analysis of V heavy chain subgroup 4

				1	Fram	ewo	ork l	11									<del></del>			
amino acid¹	92	77	78	79	80	81	82	۷	æ	ပ	83	84	85	98	87	88	83	90	91	92
Α												55	57			57				
В		• •	<del></del>			••••••	*******		<u> </u>		·				·			<u>.</u>		
· C			<u> </u>			•••••							•••••					<del></del>		57
D			**************************************	 !	1	******	••••••	**************************************				•••••		57						
E				•		1				•••••	********	•••••			*******					
F .			54			*******	*******		1	•••••		•••••	******	*******	••••••			•••••	•••••••	
G						*******		1		••••••		*******	••••	•••••	******	••••		••••••	••••••	
Н							*********	*******		••••••	•••••		******	********	*******	********				_
[			1			********		1	•••••••		3	•••••		•	••••				•••••	
K	3					46		2	*******	*******				*******			******			
L		3	1		55		53			2					********		1			
M.						1	1			1				*********			1	•••••		
N	54					3		3	1					***************************************			••••			
Р															***********			•••••		
Q		54			1	1														
R						2		2				1								
S			1	57		2	1	44	55		1				2				1	
T						1		4			53				55					
V							2	********		54		1					55			
W				******																
X		********					••••									·				
Y				•••••				******										57	56	
Z ·										-										
-		••••						•••••												
unknown (?)																				
not sequenced																				
sum of seq <sup>2</sup>	57	57	57	57	57	57	57	57	57	57	57	57	57	57	57	57	57	57	57	57
oomcaa³	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,				55	•••••••••••••••••••••••••••••••••••••••	53	44			53	55	57	57	55	57	55	57	56	57
mcaa'	N	Q	F	S	L	K	L	S	S	٧	T	Α	Α	D	T	Α	٧	Υ	Υ	С
rel. oomcaa⁵	95%	95%	95%	100%	%96	81%	93%	77%	<b>%96</b>	95%	93%	<b>%96</b>	100%	100%	%96	100%	%96	100%	%86	100%
pos occupied <sup>6</sup>	2	2	4	1	_		4			•	3	:	;	:	:	:			:	1

Table 6E: Analysis of V heavy chain subgroup 4

·										CDI	R III									
amino acid'	93	94	95	96	97	86	66	100	٧	8	ပ	۵	ш	ų.	g	I		_	×	101
Α	56		3	3	3	2	5	4	2	2	· 4		2	1		1	1	12		
В																				
. C					1				1											
D			6		5	5	5	4	3	2	4	3	1		1	2	1			41
E			- 6	1	1	2	1			1	3	1	2	1						
F.				4	1	1		2	3	2	2		1	1					31	
G			25	9	10	8	10	11	4	7	7	6	1	1	1	2	1	9		
Н		******	1				1						1			1				2
1				1		2	4	1	3	2	3		1						1	•
K			2	1						2	2			1						
L			2	6	7	3	5	3	2	4	1	5	3	3		1				
M				1	4		3	1		2	1			-					9	
N				3					2	1	1	5	1	1			2			
Р .				4	5	3	1	1	2	1	1	1	2	3	1	2	1			
Q					1	1		1			1	1			3					1
R		54	4	12	2	5	5	3	2	3	1	2	·		2	1				
S		1	1	4	8	8	1	. 2	5	7	4	2	1	1	1					
T		1	1	2	1	3	4	4	3	3			1	1	1					
V	1	1	4	2	2	5	4	4	7	3	1	2	1							•••••
w			1	2	1	2	2	4	5	1	1	2		2	1		3	2		•••••
X																		,		
Υ		•••••		1	4	5	3	6	4	2	3	4	8	4	8	3	5	8		2
Z																				
_						1	2	4	6	9	11	16	23	27	29	34	31	14	4	
unknown (?)						*******			*******					1			1	1	1	******
not sequenced			1	1	1	1	1	2	3	3	6	7	8	9	9	10	11	11	11	11
sum of seq <sup>2</sup>	57	57	56	56	56	56	56	55	54	54	51	50	49	48	48	47	46	46	46	46
oomcaa,	56	54	25		*********	*********	*********	11	7	9	11	16	23	27	29	34	31	14	31	41
mcaa'	Α	R	G	R	G	G	G	G	٧	-	-	-	-	-	-	-	-	-	F	D
rel. oomcaa⁵	98%	95%	45%	21%	18%	14%	18%	20%	13%	17%	22%	32%	47%	26%	%09	72%	%29	30%	%29	%68
pos occupied <sup>6</sup>	:	:	:																i	4

Table 6E: Analysis of V heavy chain subgroup 4

	. 1					Fra	mev	vork	· IV					1
an	nino acid'	102	103	104	105	106	107	108	109	110	111	112	113	sum
	Α						1			1				332
	В			********	******	••••		*******	*********					
<b></b>	С							•••••				•••••••	•••••••	113
<b> </b>	D		•••		••••									210
	E	**********									•••••	-		176
.	F				•••••	•••••		•				•		135
ļ	G			41		40	1					•••••		674
ļ	Н	1				•••••••••••••••••••••••••••••••••••••••				1		*******		45
		9					1						•••••••	282
<b></b>	K				3	••••								278
-	L	4				••••••		19						540
<b> </b>	M							9						43
ļ	N					•••••	1					********		204
	Р	3		*********	2	•••••		•••••				•••••	2	281
	Q			*******	29	••••••	•••••							334
<b> </b>	R	1			.4			1				••••••		250
	S	1			1			••••				36	33	986
	T				1		33	8		34				532
	V	12							36		36		-	488
	W		46											267
	Χ													
	Y	16											·	455
	Z													1
	-													466
un	known (?)	·												4
not	sequenced	10	11	16	17	17	20	20	21	21	21	21	22	426
Su	ım of seq²	47	46	41	40	40	37	37	36	36	36	36	35	
(	oomcaa¹	16	46	41	29	40	33	19	36	34	36	36	33	
	mcaa*	Υ	W	G	Q	G	T	L	٧	T	٧	S	S	
re	l. oomcaa <sup>s</sup>	34%	100%	100%	73%	100%	%68	51%	100%	94%	100%	100%	94%	
pos	s occupied <sup>6</sup>				6			4		3	1	1	2	

170

Table 6F: Analysis of V heavy chain subgroup 5

										-8				Fra	ame	wor	k I			•
amino acid'	-	2	က	4	သ	9	7	œ	6	01	=	12	13	14	15	16	17	81	19	20
. A					1			1	89		1			1						
В																				
· C							1													
D										2										
E	88	1			2				4	93						92				
F .																	1			
G	1							92							94					
Н																				
·																				96
Κ												94	94						77	
L		1		91		2												95		
М							400000000				3								1	
N																				
Р				1					1					94						
Q	. 3		92		1	90										3			1	
R						1						1	1		1				17	
S							92										94			
T							,													
V		90			89		*******		1		91									
. W																				
X																				
Y																				
. Z																				
_															********					
unknown (?)																				
not sequenced	5	5	5	5	4	4	4	4	2	2	2	2	2	2	2	2	2	2	1	1
sum of seq <sup>2</sup>	92	92	92	92	93	93	93	93	95	95	95	95	95	95	95	95	95	95	96	96
oomcaa <sup>3</sup>			••••••		•••••••		·····	<del></del>	······	<u>:</u>	91		**********	********	••••					
mcaa'	Ε	٧	Q	L	٧	Q	S	G	Α	Ε	٧	K	K	Р	G	Ε	S	L	K	l
rel. oomcaas	%96	%86	100%	%66	<b>%96</b>	92%	%66	%66	94%	%86	%96	%66	<b>%66</b>	<b>%66</b>	%66	92%	<b>%66</b>	100%	%08	100%
pos occupied <sup>6</sup>	:	:	:	:	:	i	:	:	:	•	•	:								1

Table 6F: Analysis of V heavy chain subgroup 5

								-			Τ	·		C	DRI		<del>.                                    </del>		Τ	
amino acid'	21	22	23	24	25	26	27	28	29	30	31	4	8			34	35	36	37	38
Α				3	2					4							8	-	1	
В							•		<u> </u>	<del></del>	<del></del>		·					<u> </u>		ļ
· C		96		-				1	÷	<del></del>	1	<u> </u>	<u> </u>	<u> </u>	-	<u> </u>			<u> </u>	
D	1	<u> </u>	÷	÷	<b>:</b>	<b>!</b>	ļ	2	<del></del>	ļ	2	÷		ļ	<u> </u>	<del></del>	1	<u> </u>	<u> </u>	
E	1		•	÷	<b></b>	2			<u> </u>	-	1	•			·······	<del></del>	<u> </u>	<u></u>	<u></u>	
F .	<b>[</b>		*******	<u> </u>	3	:	6		97	<del></del>	•	<u> </u>	·	2		ļ	<u></u>	······		•••••
G	1	<del></del>	•	92	······	93				<b></b>	1	<b>†</b>	ļ		ļ	<b></b>	72		•	•••••
' Н		••••••••••••••••••••••••••••••••••••••	•	<u> </u>	·······			ļ <u>.</u>	•	·······	1	·		4		ļ				1
	<b>[</b>	······	<u> </u>	<u> </u>	ļ				··········	4		<u> </u>	<u> </u>	<u></u>		93				
Κ.		<del></del>	89	<u></u>				1			<u></u>	<u> </u>	<u></u>	<u></u>					<b></b>	•••••
L		<u></u>	<u></u>	<u> </u>		••••••						<del> </del>	: :	<b></b>	1		•••••	•••••	2	•••••
М	<b> </b>		1			•••••						<u></u>	- -			1	•	•••••	1	••••••
N			1			••••••	•••••	2		4	14	<u> </u>		2		······				
Р		***********			1	********	••••••	•••••	*******	•	********		••••••							1
Q			4		•••••	•••••••	********			•••••	*******		••••	••••••	•••••	•••••				
R			1		······	1	******	2	••••••				••••••		1			******	•••••	95
S	94		••••••	1	90		•••••	84		10	61		*******	2	2		15	*******		
T	2	•••••	*********	•••••	*******	•••••	********	5			16	•••••	•••••••			2		********		
V		•••••				•••••							••••	•••••		1	•••••••••••••••••••••••••••••••••••••••	•••••	93	
W		· · ·	••••••		*********	•••••		•••••			••••••				93			97	· <u>†</u>	
X		••••••	*******		••••••	*********	**********		•••••	***************************************		••••••	•••••	*******						
Υ		•••••			•••••	•••••	90	••••••			*********			87	********					
Z				**********	•••••••		********			*********		••••••			•••••••					
												97	97							_
unknown (?)							**********				************		••••••		••••••					
not sequenced	1	1	1	1	1	1	1								*******					
sum of seq <sup>2</sup>	96	96	96	96	96	96	96	97	97	97	97	97	97	97	97	97	97	97	97	97
	94		:	:	:	:	:	:	•	•						**********			********	*******
mcaa'	S	:	K	•	•	•	Υ		********	T	S	•••••••••••••••••••••••••••••••••••••••	-	Υ	W	ı	G	W	******	R
rel. oomcaas	%86	100%	93%	%96	94%	97%	94%	2%	<b>%</b> 00	%,	%	%0(	<b>%0</b> (	%(	%	%96	%	100%	<b>%96</b>	%
nos occupiedi			••••••					ω	7	77	9	2	2	90	96	96	74			:
pos occupied <sup>a</sup>	2		5	3	4	3	.2	7	1	5	8	1	1	5	4	4	5	1	4	3

Table 6F: Analysis of V heavy chain subgroup 5

				Fr	ame	wor	k II													
amino acid'	39	40	41	42	43	44	45	46	47	48	49	20	51	52	۷	8	ပ	53	54	55
Α		•	1			1									1			2	1	
В													• • • • • • • • • • • • • • • • • • •							
· C												•		1				1		
D									<u> </u>	*******				14	•			8	93	
E					3	*********	********	97		••••					••••••		,		2	
F		*****				••••••	********					1	•••••••	2	•••••••					
G		•••••		97		96				•••••	95	********	••••••••			••••••	••••	69	1	••••••
Н			-			•••••••		······	•		•••••••			3	1	•••••		•••••		
			********			••••••		•••••		1		75	92			••••••			•••••	••••••
κ		1	•••••		94	*********		••••••							••••	••••		•••••	•••••	
L				•••••	•	••••••	94	••••••		2		2	1	*******	•••••					
М		92		*********	•••••	•	•••••	••••••		89		••••	1			*****				•••••
N		•		*******	*******	•••••	•••••	•	******				******	*******		*******	.,			
Р			96		•••••	*******	2					•••••	••••••	1	93					1
Ω	97	•••••		••••••	••••••		1					•••••		•		••••••				
R		1	****	**********	********	*******	•••••	*******			1	14		•••••••••••••••••••••••••••••••••••••••		********		1		••••••
S									•••••			1	••••••	••••••	1	·		16		96
Т		1		•••••	•••••	•••••				-		3	1		1					
V		2		••••••			••••••			5	1		••••••	••••••						
W		•••••				•••••			94	•••••••••••••••••••••••••••••••••••••••										
X				•••••	•••••	••••••	********	*******			********							<u>-</u>		
Y		••••••				••••••	*******		3					76						
Z				•••••		••••••	********	•••••												
_																97	97			
unknown (?)					*******	********	********	••••••	•••••				*******		••••••					
not sequenced					*********	•••••	••••••	•••••								******				
sum of seq'	_	97	97	97	97	97	97	97	97	97	97	97	97	97	97	97	97	97	97	97
oomcaa,	: :					:					95			•••••••••••••••••••••••••••••••••••••••			******			
mcaa*		М		G		G	L		********	М		ı	ı	Υ			-	G	D	
rel. oomcaa <sup>s</sup>	100%	95%	<b>%66</b>	100%	92%	%66	97%	%001	%/(	32%	%86	77%	92%	78%	<b>%96</b>	100%	,000 100%	71%	%9,	<b>%66</b>
pos occupied <sup>a</sup>	•					:	3	: :	: :		3									<u>ი</u> 2

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Table 6F: Analysis of V heavy chain subgroup 5

		DR	[]																<del></del>	
amino acid'	26	22	58	23	99	6	62	63	64	65	99	29	89	69	70	71	72	73	74	75
Α		6					1									88				
В																				
· C					1					1										
D	77									2							97			
E	3								2									2		
F.				2				91				1		3						
G	1									94										
Н											15									
		4	1					1				3		88						91
K			2													·		93		
L.						1		4							2					
M														3						1
N	2		14	2		********	•••••													
Р					••••	95	1		1										1	
Q				••••••	• • • • • • • • • • • • • • • • • • • •			*********	91		81	******						1		
R			78						3		1			1				1		
S	2	2			95	1	95	1					1		95	į			96	1
T		85	2		1								96							4
. V				1								93		2		9				
W																				
X																				
Y	12			92	*******															
Z -																				
_																				
unknown (?)										·····										
not sequenced																				
	•				***********	••••••	••••••		··········		:	•••••••	•••••••••••••••••••••••••••••••••••••••	•••••••••••••••••••••••••••••••••••••••	••••••••	97	•••••••••		······÷	•••••••••••••••••••••••••••••••••••••••
oomcaa,			•••••••••••		•••••	•••••		******				•••••••••••••••••••••••••••••••••••••••	96	88	•••••••••••••••••••••••••••••••••••••••	88			••••••••••	•••••••••••••••••••••••••••••••••••••••
mcaa'	D	T	R	Υ	S <sub>.</sub>	Р	S	F	Q	G	Q	٧	T		S	Α	D	K	S	1
rel. oomcaa <sup>s</sup>	79%	%88	%08	95%	%86	%86	%86	94%	94%	92%	84%	%96	%66	91%	%86	91%	100%	%96	%66	94%
pos occupied <sup>6</sup>	6	4	5	4	3		3	:	•	•			:	:	:	:	:			

Table 6F: Analysis of V heavy chain subgroup 5

		-								-			-							
				F	ram	ewo	rk I	11												
amino acid¹	9/	77	78	79	8	8	82	⋖	ဆ	ပ	83	84	82	98	87	88	68	96	91	92
Α		1	91								1	96				93				
В																				
· C							1													95
D				1										96						
E						1					1					·				
F.				1							·							2	6	
G		,						3	1							4				
Н						3														
1															2		9			
Κ											91						1			
Ĺ					96					97							2			
M																	84			
N	7							2	2						2	••••••	••••••			*********
Р			1																••••••	••••••
Q						93														
R	1						1	1	3		3									
S	87	2	1	1				90	91				96		5					
Ţ	2	94	2					1			1	1	1	·	88		1			
V			2		1	*******								1						
W							95													
χ							••••													
Υ				94			•											94	89	
Z																				
-						••••								*********						
unknown (?)						********									***********					
not sequenced																		1	2	2
sum of seq <sup>2</sup>	97	97	97	97	97	97	97	97	97	97	97	97	97	97	97	97	97	96	95	95
oomcaa³	87		91	**********	96	93	*********	***********	*********		91	•••••••••••••••••••••••••••••••••••••••	96	96	88	93	84	94	89	95
mcaa*	S	T	Α	Υ	L	Q	W	S	S	L	K	Α	S	. D	T	Α	М	Υ	Υ	С
rel. oomcaas	%06	97%	94%	97%	%66	%96	%86	93%	94%	100%	94%	%66	. %66	<b>%66</b>	91%	<b>%96</b>	87%	%86	94%	100%
pos occupied <sup>6</sup>			5		2						5				4					

Table 6F: Analysis of V heavy chain subgroup 5

			<del>-</del>							CD	RIII									
amino acid'	93	94	92	96	97	86	66	100	∢	ω	ပ	0	ш	ட	ပ	I	_	٦	×	101
Α	92		1	1	2		3	4	3	2		1			1			4		2
В																				
· c						1	1	1			2		1		••••••					
D				3	3	3	3	1	2	1	1	2		2	1	1	2			37
E			1	1	1	2	*********		1	1				1			1		*******	
F .					1		3			3	2		1	•	*********		••••••••••••••••••••••••••••••••••••••		26	
G			1	9	11	12	12	5	2	4	3.	. 10	2	1	•••••	••••••	)	5	•••••	
Н			10	1		2			1	1		1								
l				3		2	2	1	1	4	1	1	·	1	1	••••••				
K		1	1	1		1	3	1								2				
L			11	2	3	1	1	2	5		1		1		1					
M					2	1	1		1	1	1	1							10	
N				1		2		1	1	2			1				••••	2		
P ·			5	1	4	3	1	2				1	1	. 1	1					
Q		1	3	2		1	1	4	2	1	2									3
R		92	7	9	2	2		2	1		2									
S		1	1	3	2	6	4	4	5	3	5	3	2	2			1		1	
T	1		1	3	2	1	2	6	3	3	6	1		1			-			
· V	2		2	4	4		1		1	2			1							
W			1		2	1					1		2		1		1	1		
. X																				
Υ				1	6	3	6	9	8	7	2	1	2	6	8	9	9	10	***************************************	1
Z												-				********				
-						1	1	2	8	10	16	23	30	30	31	32	30	22	7	2
unknown (?)													1		***************************************	1	1	1	***************************************	***************************************
not sequenced	2	2	52	52	52	52	52	52	52	52	52	52	52	52	52	52	52	52	53	52
sum of seq <sup>2</sup>	95	95	45	45	45	45	45	45	45	45	45	45	45	45	45	45	45	45	44	45
oomcaa,	92	92	11	9	11	12	12	9	8	10	16	23	30	30	31	32	30	22	26	37
mcaa⁴	Α	R	L	G	G	G	G	Υ	Υ	-	-	-	-	-	-	-	-	-	F	D
rel. oomcaa <sup>s</sup>	92%	97%	24%	20%	24%	27%	27%	20%	18%	22%	36%	51%	%29	9/2	%69	71%	%29	49%	29%	82%
pos occupied <sup>a</sup>				•	:		•	•	:	:				:	:					5

Table 6F: Analysis of V heavy chain subgroup 5

				•	Fra	mev	work	· IV	<del></del> -				Ì
amino acid'	102	103	104	105	106	107	108	109	110	111	112	113	sum
А												1	611
В											•	••••••	
С												******	205
D	1	*************	********	*******		•	********	*********	•••••	•••••	•		458
E		***************************************	***************************************	1				*******	*********				404
F	2	*******			***************************************					•••••	•••••		256
G		************	41		41	•		••••••	•••••		•		1065
Н		***********	**********			•••			•••••	••••••			44
1	9		***********			•	•		2	•••••			588
K				3								********	650
L	2						25	1					549
М							8						303
N													64
Р	2					1					1		414
Q				34									612
R				3									351
S	2										40	39	1545
T	1					40	8		39				604
V	11							40		41			594
W		43	*************	,									432
X													
Υ	13		•••••										738
Z			-										
	2												635
unknown (?)				•	.,			•••••		••••			4
not sequenced	52	54	56	56	56	56	56	56	56	56	56	57	1678
sum of seq <sup>2</sup>	45	43	41	41	41	41	41	41	41	41	41	40	
oomcaa,	13	*************	*********	*******	• • • • • • • • • • • • • • • • • • • •	• • • • • • • • • • • • • • • • • • • •	*********	40		********			
mcaa <sup>4</sup>	Υ	W	G	Q	G	T	L	٧	T	٧	S	S	
rel. oomcaas	29%	100%	100%	83%	100%	%86	61%	98%	95%	100%	98%	%86	
pos occupied <sup>6</sup>	:	:		:	:	•	:			1	2	2	

Table 6G: Analysis of V heavy chain subgroup 6

•														Fr	ame	wor	kΙ			
amino acid'	_	7	က	4	2	9	7	æ	6	10	=	12	13	14	15	16	17	18	19	20
A												1								
В														·						
· c					<del> </del>			:	********	*********							*********	***********	*******	
D				<u> </u>	<del></del>	·						•••••	•••••	••••••			•••••			
E	**********					······································			••••	*******	•••••	_	•••••	********		••••••	••••	*********	*****	•••••••
F .		••••••					********		********	•••••	•••••	•••••	••••••		••••••		•••••	•••••		
G		•				••••••	••••	52	•••••••	67	••••••		********	••••••		**********		********	********	••••••
. Н		••••••			•••••	••••••	•		••••••	*******			•••••	•••••	********			•••••	••••••	*********
1	*******	•••••••	•••••		•••••••		********	•••••		••••••	•••••		******	*******	•••••	••••••	••••••	•••••		********
K			********			••••	*******	••••					68	*******		••••••				
L		•		52			•	••••	••••••	•••••	68	1	*******	••••••	••••			67	1	68
М			•••••	•••••	•••••	••••		•••	••••••					••••••	••••	••••••		***************************************		•••••
N				•••••	••••••	•••••••		•••••	••••				••••••		•	••••••			*********	
Р			••••••		•	*********	*********	•	68		•••••		••••••	67	•••••		•••••		1	•••
Q	52	••••	52	•••••	51	52	**********	*****		*******				*********	*******	68		•••••••		
R					1	-	********			1				*********						
S				••••••	******	••••	52			********			•••••••••••••••••••••••••••••••••••••••	1	68			•••••••	66	100000
· T					•••••	******	*******			***************************************					••••••		68			
V		52					******			********		66		••••••	*******			1		***************************************
W			******	*******		********	•••••	••••••	********	*********								******	******	
X	*********		•		********	••••••	**********	•••••	***************************************				*******		•••••	***********				***************************************
Υ							*********		***************************************	***************************************					••••••	***************************************			*******	
Z				•	••••••	*******	*********		***************************************	********	••••••		•	•••••	•••••					
-																				
unknown (?)										•••••			•••••••••••••••••••••••••••••••••••••••							
not sequenced	22	22	22	22	22	22	22	22	6	6	6	6	6	6	6	6	6	6	6	6
sum of seq <sup>2</sup>	52	52	52	52	52	52	52	52	68	68	68	68	68	68	68	68	68	68	68	68
oomcaa¹									:		68	•••••••		••••••	***********	************				
mcaa•	Q	٧	Q	L	Q	Q	S	G	Ρ	G	L	٧	K	Ρ	S	Q	T	L	S	L
rel. oomcaa⁵	100%	100%	100%	100%	%86	100%	100%	100%	100%	%66	100%	9/0/6	100%	%66	100%	100%	%00 <sub>1</sub>	%66	37%	100%
pos occupied <sup>6</sup>	1	1	1	1				:	:	:			•••••••••••••••••••••••••••••••••••••••	•••••••••••••••••••••••••••••••••••••••				•	•	

Table 6G: Analysis of V heavy chain subgroup 6

												•		CC	RI					
amino acid'	21	22	23	24	25	26	27	28	29	30	31	٧	æ	32	33	34	35	36	37	38
A	1		67											66	67					
В																				
С		68						***********							******					
D							68				1						1			
E																				
F .										2				1	1				1	
G			1			69							3	1	2					
Н																	1			
				64								2					1		70	
K												3								
L																				
М					••••••				*******						*********					
N					•••••	•••••	1				2	66				*******	70			******
Р						••••••	•••••						•••••		**********	•				
Q			•		*********	•••••	•••••	••••••	•••••	•••••		•••••			••••••	**********				
R				•••••	*******	********	*********		********	•	2	1			********	••••		•		74
S	1		*******	1	69	•••••	•••••	69		68	66	*********	67	•••••	3		1			
T	67					*******	*****		*******		2	1	4	******	1			•••••	•••	
V			1	4		•••••	••••		70			•••••	•••••	6	******	••••••			2	
W		1		•••••	********	•••••	•				•••••	•••••		•••••	********	74		74		
Х					*******	••••••								•••••••						
Y		••••••	••••••	**********	*******	*********			• • • • • • • • • • • • • • • • • • • •			1		*******	•••••	*********		•••••	1	
Z	1	•	•••••	•••••			•••••	*********		**********		-	•••••	•••••	*******		••••••••••••••••••••••••••••••••••••••		•••••	
_																				
unknown (?)							••••••	*******	****		1		•••••	••••••	••••••	*********				
not sequenced	5	5	5	5	5	5	5	5	4	4								: :		
sum of seq <sup>2</sup>	69	69	69	69	69	69	69	69	70	70	74	74	74	74	74	74	74	74	74	74
oomcaa³	67	68	67	64	69	69	68	69	70	68	66	66	67	66	67	74	70	74	70	74
mcaa*	T	С	Α	1	S	G	D	S	٧	S	S	N	S	Α	Α	W	N	W	ī	R
rel. oomcaa <sup>s</sup>	92%	%66	92%	93%	100%	100%	%66	100%	100%	%/£	%68	39%	31%	39%	91%	100%	95%	100%	92%	100%
pos occupied <sup>a</sup>		7	:	:	:	<u> </u>	:	:	:	:	:	:	:	:	:	:	:	:	:	<b>†</b>

Table 6G: Analysis of V heavy chain subgroup 6

				Fr	ame	wor	k II						Π							
amino acid'	39	40	41	42	43	44	45	46	47	48	49	20	51	52	۷	8	ပ	53	54	55
Α				1									1					1		
В		:																		
· C	<u> </u>	<u> </u>														:				
D			<u> </u>																	
E	<b>.</b>	<u></u>	į	<u>.</u>				74							·					
F .	<u> </u>	<u></u>	ļ	<u> </u>										2	1			1		
G	<b></b>	<b>.</b>	ļ	<u>.</u>		74					74	1							1	
Н	ļ		<u> </u>	<u></u>	<u>.</u>										1					
·	<b>.</b>		<u></u>	<u>.</u>	<u>.</u>															
K	1		<u> </u>	<u></u>	1											1			66	
<u> </u>	1		<u> </u>				74			74										
M							********													
N					•••••			•••••											1	
Р		•••••	73				*******	********												
Q	72	*********	ļ	<b>6</b>			•••••													
R					73							73				72			1	1
S		74	1	73						·····i						1		72		
Т													73						5	
V																				
W		•••••							74											73
X		•••••																		
Υ .		··	•••••											72	72					
<u>Z</u>								-												
																	74			
unknown (?)																				
not sequenced		-																		
sum of seq'	74	74	74	74	74	74	74	74	74	74	74	74	74	74	74	74	74	74	74	74
oomcaa,	: :			73		74	74	74	74	74	74	73	73	72	72	72	74	72	66	73
mcaa'	Q	S	Р	S	R	G	L	Ε	W	L	G	R	T	Υ	Υ	R	-	S	K	W
rel. oomcaas	92%	100%	<b>%66</b>	%66	%66	100%	100%	100%	100%	100%	100%	%66	%66	%26	92%	92%	100%	92%	%68	%66
pos occupied <sup>6</sup>	:	i	_ :	•	2	1	1		1 88	1		•	2	:			·····			:

Table 6G: Analysis of V heavy chain subgroup 6

	С	DR	11		-															
amino acid'	26	57	28	59	9	61	62	63	64	65	99	67	89	69	70	71	72	73	74	75
Α					73	1							2			6		1		
В							P 0 0 0 1 1 1 1 1			•	•••••		•••••		*********	•	••••	•••••		
С				1									***************************************	•						
D		***************************************	68			1	********		••••					•••••••	2		73			
E	1		3			7	******		1						••••••	•••••				2
F.	7			*******	******	•	*******					•		**********		•	.,,,			
G			1				1			8			•••••	************		••••••	•			
Н	1													•••••••			1			
1						1						65	2	71				1		
K		1							67						1					70
L	1					5		2				4						1		
M												1								
N	2	65	1						1						69					
Р					1	1				,						66				
Q ·									2		· 1									
R		1							3		73									
S .	2	2	1	1			73			66			1		2	1			73	
T		4											69	1				71	1	2
V					******	58		72				4		2		1				
W	.,		*****				•													
Χ							•••••													
Υ	60	1	•••••	72	********	*******			******					**********						
Z																				
-				<u>.</u>			•	••••••												
unknown (?)				<u>.</u>					•••••											
not sequenced																				
sum of seq <sup>2</sup>	74	74	74	74	74	74	74	74	74	74	74	74	74	74	74	74	74	74	74	74
oomcaa¹	60	65	68	72	73	58	73	72	67	66	73	65	69	71	69	66	73	71	73	70
mcaa'	Υ	N	D	Υ	Α	٧	S	٧	K	S	R	١	T	1	N	Р	D	T	S	K
rel. oomcaas	81%	%88	92%	92%	%66	78%	%66	97%	91%	%68	%66	988%	93%	<b>%96</b>	93%	%68	<b>%66</b>	<b>%96</b>	%66	95%
pos occupied <sup>6</sup>	7	6	5	:	:	:	:	:	:	:			4	: :			2			3

Table 6G: Analysis of V heavy chain subgroup 6

					Fran	new	ork l	11												
amino acid'	9/	77	78	79	80	81	82	∢	8	υ	83	84	85	98	87	88	83	90	91	92
Α													1			74				
В													<u></u>				<u> </u>			<del></del>
· c							<u> </u>				<u></u>	<u></u>				<u> </u>	<del></del>	<del></del>	<u> </u>	73
D				<u> </u>				3			·····			73		<del></del>		<u> </u>	<del></del>	<del></del>
E		<del></del>	T		<b>!</b>				······				73		<del> </del>	<del></del>	<del>.</del>	<del> </del> -	<del>:</del>	
F			71				†	······································	1					*******			·······	<del> </del>	3	
G										•••••				1				•		
Н						2		1	•	•••••	•	<b></b>	••••	••••••	}	•	••••••			-
			1			••••••	********	********	,				*********	*******			2			
K								4		•••••	•••••	*****	*******		••••••		•••••			
L	·	1			74		72				*********	•••••	*******	•••••	********					
М							1			1			************		********	•••••	2			
N	74							63			•••••				********	•	*******	••••••	1	
Р		****										70			*******			••••••		*******
Q		72				71											••••	•	****	
R		1				1		1									**********	••••••		1
S				74	****	••••		1	73		1	3			******	*********		•		
Т			******					1			73			•••••••	74			1		
V			2				1			73							70			
W						***************************************														*******
X		*********															•••••			
Y																		73	70	••••••
Z																				
-																				
unknown (?)																				
not sequenced												1								
sum of seq²	74	74	74	74	74	74	74	74	74	74	74	73	74	74	74	74	74	74	74	74
oomcaa <sup>3</sup>				74	74	71	72	63	73	73	73	70	73	73	74	74	70	73	70	73
mcaa'	N	Q	F	S	L	Q	L	N	S	٧	T	Р	Е	D	T	Α	٧	Υ	Υ	С
rel. oomcaa <sup>s</sup>	100%	92%	%96	100%	100%	%96	92%	85%	%66	%66	%66	%96	%66	%66	100%	100%	95%	%66	95%	<b>%66</b>
pos occupied <sup>6</sup>	i	i	3	1		•	•		:	:	•	•		*******		1		********		

Table 6G: Analysis of V heavy chain subgroup 6

	CDR III																			
amino acid'	93	94	95	96	97	98	99	100	⋖	80	U	۵	ш	ய		Ξ	_	_	×	101
Α	69		11	1	3	12	4	3	2	5		8						10	1	
В											•		•••••	*******	** ** ***					
· C					1		1			1		1	1							
D			19	4	3	7	4	3	1	6	1	1	1	0- 000 00						62
E			10	4	2	1	2	2	1	2			••••••	••••••	•••••		1			
F .	1		1	1	1		1	2	3	•	2	••••	•	1	*******	**********			38	4
G	1		16	4	15	15	11	8	6	2	5	1	8	6	1			17		
Н				1		1			1	1	1	1				1	1	1		
1				1	2		2		5	1										
Κ		1	1	1	1	1	1	1				1						<u> </u>		
L		******	1	8	4	2	3	2	1					1	5			<u> </u>	8	
M				1				1			5								11	
N			1	3	1	2	1	1	1	3		2		1		1	3			
Р				10	4		5	3		5	1		1							
Q			1	1	1	1					1									1
R		69	1	7	8	1	8	8	3		1	1	5							1
S		3	5	5	5	7	6	7	3	4	2					1	1			
T			1	1	4	3	4	4	6	3	1			1				<u>i</u>		
V	3	1	4	5	1	9			4		9	5	1	1		·			2	
W			1	6	8		3	2	4								4	4		
X							****													
Y				6	4	2	2	2	6	6	2	4	2	1	8	8	12	12		
Z						,														
				2	3	7	14	23	25	33	41	47	53	54	57	56	50	28	12	4
unknown (?)							•••••							6	1	5		<u> </u>		
not sequenced				1	2	2	1	1	1	1	1	1	1	1	1	1	1	1	1	1
sum of seq <sup>2</sup>	74	74	73	72	71	71	72	72	72	72	72	72	72	72	72	72	72	72	72	72
oomcaa <sub>3</sub>	69	69	19	10	15	15	14	23	25	33	41	47	53	54	57	56	50	·28	38	62
mcaa'	Α	R	D	Р	G	G	-	-	-	-	-	-	-	-	-	-	-	-	F	D
rel. oomcaas	93%	93%	26%	14%	21%	21%	19%	32%	35%	46%	57%	65%	74%	75%	%62	78%	%69	39%	53%	%98
pos occupied <sup>6</sup>	•		•					: :	: :			:								5

Table 6G: Analysis of V heavy chain subgroup 6

	•		Framework IV											
	amino acid'	102	103	104	105	106	107	108	109	110	Ξ	112	113	sum
	Α							2						494
	В				·····							<del></del>	<del></del>	
	С				<del> </del>	<del></del>	<del> </del>	<u> </u>	<del></del>			<del></del>	<del></del> -	147
	D					<del></del>			1	<del> </del>		<del></del>		403
	E					<del></del>	······				·	<del></del>		186
	F	2			······	<del></del>	••••••••••••••••••••••••••••••••••••••	ļ				2	•	150
:	G			49	••••••••••••••••••••••••••••••••••••••	50		<b>†</b>				······		571
	Н	2				•	••••••	·······	•					18
		9		}			3	<b></b>	1					304
	K		······		1			1						293
	L	5	•		**********		*******	26					********	632
	M				••••••		*******	8	••••••		-	-		31
İ	N												********	436
	Р	4			6						*******		1	387
	Q				40									539
	R				2									495
	S	4		1			1					43	46	1271
	Ţ						45	4		45	·			640
	<u> </u>	21						2	46		48			647
	W		65					5						398
	X		******		********		*******							
	Υ	19	~					••••••			•••••			518
	Z													
		2								******	*********			585
l	unknown (?)													13
	not sequenced	5	8	23	24	23	24	25	25	28	25	28	26	580
	sum of seq <sup>2</sup>	68	65	50	49	50	49	48	48	45	48	45	47	
	oomcaa <sub>a</sub>	21	********		************				46	45	48	43	46	
	mcaa⁴	٧	W	G	Q	G	T	L	٧	T	٧	S	S	
	rel. oomcaas	31%	100%	%86	82%	100%	92%	54%	%96	100%	100%	%96	%86	
	pos occupied <sup>6</sup>	:	:	:	:	:		•		1	1	2		

# Appendix to Tables 1A-C

#### A. References of rearranged sequences

### References of rearranged human kappa sequences used for alignment

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## Claims

1. A method of setting up one or more nucleic acid sequences encoding one or more (poly)peptide sequences suitable for the creation of libraries of (poly)peptides said (poly)peptide sequences comprising amino acid consensus sequences, said method comprising the following steps:

- (a) deducing from a collection of at least three homologous proteins one or more (poly)peptide sequences comprising at least one amino acid consensus sequence;
- (b) optionally, identifying amino acids in said (poly)peptide sequences to be modified so as to remove unfavorable interactions between amino acids within or between said or other (poly)peptide sequences;
- (c) identifying at least one structural sub-element within each of said (poly)peptide sequences;
- (d) backtranslating each of said (poly)peptide sequences into a corresponding coding nucleic acid sequence;
- (e) setting up cleavage sites in regions adjacent to or between the ends of sub-sequences encoding said sub-elements, each of said cleavage sites:
  - (ea) being unique within each of said coding nucleic acid sequences;
  - (eb) being common to the corresponding sub-sequences of any said coding nucleic acids.
- 2. A method of setting up two or more sets of one or more nucleic acid sequences comprising executing the steps described in claim 1 for each of said sets with the additional provision that said cleavage sites are unique between said sets.
- 3. The method of claim 2 in which at least two of said sets are deduced from the same collection of at least three homologous proteins.
- 4. The method according to any one of claims 1 to 3, wherein said setting up further comprises the synthesis of said nucleic acid coding sequences.
- 5. The method according to any one of claims 1 to 4, further comprising the cloning of said nucleic acid coding sequences into a vector.

6. The method according to any one of claims 1 to 5, wherein said removal of unfavorable interactions results in enhanced expression of said (poly)peptides.

- 7. The method according to any one of claims 1 to 6, further comprising the steps of:
  - (f) cleaving at least two of said cleavage sites located in regions adjacent to or between the ends of said sub-sequences; and
  - (g) exchanging said sub-sequences by different sequences; and
  - (h) optionally, repeating steps (f) and (g) one or more times.
- 8. The method according to claim 7, wherein said different sequences are selected from the group of different sub-sequences encoding the same or different sub-elements derived from the same or different (poly)peptides.
- 9. The method according to claims 7 or 8, wherein said different sequences are selected from the group of:
  - (i) genomic sequences or sequences derived from genomic sequences;
  - (ii) rearranged genomic sequences or sequences derived from rearranged genomic sequences; and
  - (iii) random sequences.
- 10. The method according to any one of claims 1 to 9 further comprising the expression of said nucleic acid coding sequences.
- 11. The method according to any one of claims 1 to 10 further comprising the steps of:
  - (i) screening, after expression, the resultant (poly)peptides for a desired property;
  - (k) optionally, repeating steps (f) to (i) one or more times with nucleic acid sequences encoding one or more (poly)peptides obtained in step (i).
- 12. The method according to claim 11, wherein said desired property is selected from the group of optimized affinity or specificity for a target molecule, optimized enzymatic activity, optimized expression yields, optimized stability and optimized solubility.

13. The method according to any one of claims 1 to 12, wherein said cleavage sites are sites cleaved by restriction enzymes.

- 14. The method according to any one of claims 1 to 13, wherein said structural sub-elements comprise between 1 and 150 amino acids.
- 15. The method according to claim 14, wherein said structural sub-elements comprise between 3 and 25 amino acids.
- 16. The method according to any one of claims 1 to 15, wherein said nucleic acid is DNA.
- 17. The method according to any one of claims 1 to 16, wherein said (poly)peptides have an amino acid pattern characteristic of a particular species.
- 18. The method according to claim 17, wherein said species is human.
- 19. The method according to any one of claims 1 to 18, wherein said (poly)peptides are at least part of members or derivatives of the immunoglobulin superfamily.
- 20. The method according to claim 19, wherein said members or derivatives of the immunoglobulin superfamily are members or derivatives of the immunoglobulin family.
- 21. The method according to claim 19 or 20, wherein said (poly)peptides are or are derived from heavy or light chain variable regions wherein said structural sub-elements are framework regions (FR) 1, 2, 3, or 4 or complementary determining regions (CDR) 1, 2, or 3.
- 22. The method according to claim 20 or 21, wherein said (poly)peptides are or are derived from the HuCAL consensus genes: Vκ1, Vκ2, Vκ3, Vκ4, Vλ1, Vλ2, Vλ3, VH1A, VH1B, VH2, VH3, VH4, VH5, VH6, Cκ, Cλ, CH1 or any combination of said HuCAL consensus genes.
- 23. The method according to any one of claims 20 to 22, wherein said derivative of said immunoglobulin family or said combination is an Fv, disulphide-linked Fv, single-chain Fv (scFv), or Fab fragment.

The method according to claims 22 to 23, wherein said derivative is an scFv fragment comprising the combination of HuCAL VH3 and HuCAL Vλ2 consensus genes that comprises a random sub-sequence encoding the heavy chain CDR3 sub-element.

- 25. The method according to any one of claims 1 to 24, wherein at least part of said (poly)peptide sequences or (poly)peptides is connected to a sequence encoding at least one additional moiety or to at least one additional moiety, respectively.
- 26. The method according to claim 25, wherein said connection is formed via a contiguous nucleic acid sequence or amino acid sequence, respectively.
- 27 The method according to claims 25 to 26, wherein said additional moiety is a toxin, a cytokine, a reporter enzyme, a moiety being capable of binding a metal ion, a peptide, a tag suitable for detection and/or purification, or a homo- or hetero-association domain.
- 28. The method according to any one of claims 10 to 27, wherein the expression of said nucleic acid sequences results in the generation of a repertoire of biological activities and/or specificities, preferably in the generation of a repertoire based on a universal framework.
- 29. A nucleic acid sequence obtainable by the method according to any of claims 1 to 28.
- 30. A collection of nucleic acid sequences obtainable by the method according to any of claims 1 to 28.
- 31. A recombinant vector obtainable by the method according to any of claims 5 to 28.
- 32. A collection of recombinant vectors obtainable by the method according to any of claims 5 to 30.
- 33. A host cell transformed with the recombinant vector according to claim 31.

34. A collection of host cells transformed with the collection of recombinant vectors according to claim 32.

- 35. A method of producing a (poly)peptide or a collection of (poly)peptides as defined in any of claims 1 to 28 comprising culturing the host cell according to claim 33 or the collection of host cells according to claim 34 under suitable conditions and isolating said (poly)peptide or said collection of (poly)peptides.
- 36. A (poly)peptide devisable by the method according to any one of claims 1 to 3, encoded by the nucleic acid sequence according to claim 29 or obtainable by the method according to any one of claims 4 to 28 or 35.
- 37. A collection of (poly)peptides devisable by the method according to any one of claims 1 to 3, encoded by the collection of nucleic acid sequences according to claim 30 or obtainable by the method according to any one of claims 4 to 28 or 35.
- 38. A vector suitable for use in the method according to any of claims 5 to 28 and 35 characterized in that said vector is essentially devoid of any cleavage site as defined in claim 1(e) and 2.
- 39. The vector according to claim 38 which is an expression vector.
- 40. A kit comprising at least one of;
  - (a) a nucleic acid sequence according to claim 29;
    - (b) a collection of nucleic acid sequences according to claim 30;
    - (c) a recombinant vector according to claim 31;
    - (d) a collection of recombinant vectors according to claim 32;
    - (e) a (poly)peptide according to claim 36;
    - (f) a collection of (poly)peptides according to claim 37;
    - (g) a vector according to claim 38 or 39; and optionally,
    - (h) a suitable host cell for carrying out the method according to claim 35.
- 41. A method of designing two or more genes encoding a collection of two or more proteins, comprising the steps of:

- (a) either
  - (aa) identifying two or more homologous gene sequences, or
  - (ab) analyzing at least three homologous genes, and deducing two or more consensus gene sequences therefrom,
- (b) optionally, modifying codons in said consensus gene sequences to remove unfavourable interactions between amino acids in the resulting proteins,
- (c) identifying sub-sequences which encode structural subelements in said consensus gene sequences
- (d) modifying one or more bases in regions adjacent to or between the ends of said sub-sequences to define one or more cleavage sites, each of which:
  - (da) are unique within each consensus gene sequence,
  - (db) do not form compatible sites with respect to any single sub-sequence,
  - (dc) are common to all homologous sub-sequences.
- **42.** A method of preparing two or more genes encoding a collection of two or more proteins, comprising the steps of :
  - (a) designing said genes according to claim 41, and
  - (b) synthesizing said genes.
- 43. A collection of genes prepared according to the method of claim 42.
- 44. A collection of two or more genes derived from gene sequences which:
  - (a) are either homologous, or represent consensus gene sequences derived from at least three homologous genes, and

- (b) carry cleavage sites, each of which:
  - (ba) lie at or adjacent to the ends of genetic sub-sequences which encode structural sub-elements,
  - (bb) are unique within each gene sequence,
  - (bc) do not form compatible sites with respect to any single subsequence, and
  - (bd) are common to all homologous sub-sequences.
- 45. The collection of genes according to either of claims 43 or 44 in which each of said gene sequences has a nucleotide composition characteristic of a particular species.
- 46. The collection of genes according to claim 45 in which said species is human.
- 47. The collection of genes according to any of claims 43 to 46 in which one or more of said gene sequences encodes at least part of a member of the immunoglobulin superfamily, preferably of the immunoglobulin family.
- 48. The collection of genes according to claim 47 in which said structural subelements correspond to any combination of framework regions 1, 2, 3, and 4, and/or CDR regions 1, 2, and 3 of antibody heavy chains.
- 49. The collection of genes according to claim 47 in which said structural subelements correspond to any combination of framework regions 1, 2, 3, and 4, and/or CDR regions 1, 2, and 3 of antibody light chains.
- **50**. A collection of vectors comprising a collection of gene sequences according to any of claims 43 to 49.

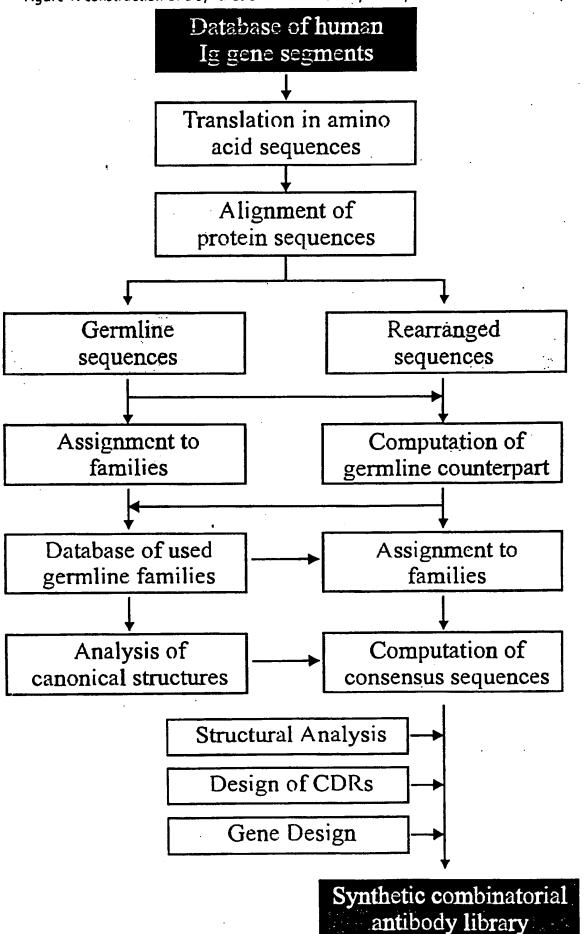
51. The collection of vectors according to claim 50 comprising the additional feature that the vector does not comprise any cleavage site that is contained in the collection of genes according to any of claims 43 to 49.

- 52. A method for identifying one or more genes encoding one or more proteins having a desirable property, comprising the steps of:
  - (a) expressing from the collection of vectors according to either of claims 50 or 51 a collection of proteins.
  - (b) screening said collection to isolate one or more proteins having a desired property,
  - (c) identifying the genes encoding the proteins isolated in step (b),
  - (d) optionally, excising from the genes encoding the proteins isolated in step (b) one or more genetic sub-sequences encoding structural subelements, and replacing said sub-sequence(s) by one or more second sub-sequences encoding structural sub-elements, to generate new vectors according to either of claims 50 or 51,
  - (e) optionally, repeating steps (a) to (c).
- 53. A method for identifying one or more genes encoding one or more antibody fragments which binds to a target, comprising the steps of:
  - (a) expressing from the collection of vectors according to either of claims 50 or 51 a collection of proteins,
  - (b) screening said collection to isolate one or more antibody fragments which bind to said target,
  - (c) identifying the genes encoding the proteins isolated in step (b),
  - (d) optionally, excising from the genes encoding the antibody fragments isolated in step (b) one or more genetic sub-sequences encoding structural sub-elements, and replacing said sub-sequence(s) by one or

more second sub-sequences encoding structural sub-generate new vectors according to either of claims 50 or 51,

- (e) optionally, repeating steps (a) to (c).
- 54. A kit comprising two or more genes derived from gene sequences which:
  - (a) are either homologous, or represent consensus gene sequences derived from at least three homologous genes, and
  - (b) carry cleavage sites, each of which:
    - (ba) lie at or adjacent to the ends of genetic sub-sequences which encode structural sub-elements,
    - (bb) are unique within each gene sequence,
    - (bc) do not form compatible sites with respect to any single subsequence, and
    - (bd) are common to all homologous sub-sequences.
- 55. A kit comprising two or more genetic sub-sequences which encode structural sub-elements, which can be assembled to form genes, and which carry cleavage sites, each of which:
  - (a) lie at or adjacent to the ends of said genetic sub-sequences,
  - (b) do not form compatible sites with respect to any single sub-sequence, and
  - (d) are common to all homologous sub-sequences.

Figure 1: construction of a synthetic human antibody library based on consensus sequences



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Figure 2B: VL lambda consensus sequences

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Figure 2B: VL lambda consensus sequences

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e sequ	ACC IGG	H	rac atg	Ø	~~~~ GCAGAAACCA CGTCTTTGGT	OI .	CAA
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Figure 3A: V kappa 1 (Vk.1) gene sequence. DIQMTQ ECORV	GA	E	\CC 166	O,	.∼ ;GA		CT
/ карра О .V	~~~~~ GATATCCAGA CTATAGGTCT		TCGTGTGACC AGCACACTGG	/ Y KpnI	CGTGGTACCA GCACCATGGT	လ	CAG
igure 3A: V k D I ECORV	~~~~~ GATATC CTATAG	Α >	GTG CAC	ĭ M	TGG ACC	ഗ	CAG
可可	GA'CT		T AG	A	, , , , ,	Ø	Ü

gure 3A: V kappa 1 (Vk1) gene sequence (continued) CGGTCGTCGA ACGTTTCGCC CCAGGGCAGG GCAAAATCGC CGAGACCTAG	TLTISSLQPEDF Eco57I	~~~~~~ BbsT	<pre></pre>	GAAGACTTTG	CTTCTGAAAC	F G Q MscI	
)GC	P ECO	<b>?</b>		CJ	3GA	H	
ATC	OI			AAC	TL	വ	
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J)				CCA	3GT/	Ä	
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اد (وم 200	Ä			CTG	GAC	O.	
ranen I'T'T	H			ACC	rgg I	Q	
gene se	Ē.			TTT.	GCCGTGACTA AAATGGGACT	Ö	
Figure 3A: V kappa 1 (Vk1) gene sequence (continued) CGGTCGTCGA ACGTTTCGCC C	Ω			AT	TA	>-	
	Ω Ή 9			CTG	GAC	A T Y Y	
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CTTTGGCCAG GAAACCGGTC CCCCGCCGAC GGGCGGCTG CATTATACCA GTAATATGGT TTGCCAGCAG AACGGTCGTC CGACCTATTA GCTGGATAAT

G T K V E I K R T Bsiwi GGTACGAAAG TTGAAATTAA ACGTACG CCATGCTTTC AACTTTAATT TGCATGC

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Figure 3B: V kappa 2 (Vk2) gene sequence

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Дı	CTGCCAGTGA GACGGTCACT	. <b>ப</b>	AAGCCTGCTG TTCGGACGAC	P G SexAI	AACCAGGTCA TTGGTCCAGT
ü	TGC	W	AGO	SA	ACC
		$\mathbf{C}$	4 H	×	•
W	TGACCCAGAG CCCACTGAGC ACTGGGTCTC GGGTGACTCG	O	ATTAGCTGCA GAAGCAGCCA AAGCCTGCTG TAATCGACGT CTTCGTCGGT TTCGGACGAC	Q	TCTGGATTGG TACCTTCAAA AGACCTAACC ATGGAAGTTT
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D I EcoRV	GATATCGTGA CTATAGCACT	Д	GCCTGCGAGC CGGACGCTCG	G Y N	GCTATAACTA CGATATTGAT
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(continued)
gene sequence
2 (VK2)
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Figure 3B:

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		200 200	E+		CCA		
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	н	TT	<b>&gt;</b>		TA	R T Bsiwi	ATTAAACGTA C TAATTTGCAT G
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kappa	ტ	19CJ	A E E		GAZ		CT.
Figure 3B: V kappa 2 (Vx2) gene sequence (continued)	Ø	TAGCGGCTCT ATCGCCGAGA	A E		AAGCTGAAGA TTCGACTTCT	<b>[-</b>	CCGACCTTTG GGCTGGAAAC
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<b>-</b> -	<b>-</b>		CTGAGCCTGT	GACTCGGACA		S C R A S Q S V S	-		GAGCGTGAGC	CTCGCACTCG	H			CACCGCGTCT	GTGGCGCAGA
U	2		CCC	CGG		>			GTG	CAC	<u>m</u>			CCG	CGC
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^	C C		SCC	SCGC					GAG	CTC	ტ	H	<b>?</b>	GT(	CAC
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seque			CCC	3GG	•	ഗ		₹	AGC	rcg				GCA	CGT
Figure 3C: V kappa 3 (Vk3) gene sequence	기 기 >		GATATCGTGC TGACCCAGAG CCCGGCGACC	CTATAGCACG ACTGGGTCTC GGGCCGCTGG		ᆸ			CTGAGCTGCA · GAGCGAGCCA	GACTCGACGT CTCGCTCGGT	Ø		<b>?</b>	TGGCGTGGTA CCAGCAGAAA CCAGGTCAAG	ACCGCACCAT GGTCGTCTTT GGTCCAGTTC
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kappa		,	GT(	3CA(		~			CGA(	3CT(	3	乙	<b>?</b>	rgg'	ACC.
3C: V	D L EcoRV	~ ~ ~ ~ ~	ATC	TAC		R A T			ACGTGCGACC	TGCACGCTGG	L A W Y			3CG.	CGC
Figure	ച ല്	<b>?</b>	GA1	$CT^{\prime}$		ہلم			ACC	TG(	Ц			TG(	AC(

Figure 3C: V kappa 3 (Vk3) gene sequence (continued)

T CGCCGAGACC	P E D Eco57I	?	BbsI	CCT	P T F G MscI	GACCTT CTGGAA		
CGCGCAAAAT	S I			CAGCCTGGAA GTCGGACCTT	T T	CCACCCCGCC		
ACCCCAGGGC	EH N			TGACCATTAG CAGCCTGGAA ACTGGTAATC GTCGGACCTT	У Н	CAGCATTATA GTCGTAATAT	K R T Bsiwi	TAAACGTACG ATTTGCATGC
CGGCACGTTG ACCCCAGGGC	D F T L		•	GATTTTACCC CTAAAATGGG	Z Z	TTATTGCCAG	K V E I	CAGGGTACGA AAGTTGAAAT
CCGCGCTCGT	S D		BamHI	ATCCGGCACG TAGGCCGTGC	F A V Y	TTGCGGTGTA	G T CI	CAGGGTACGA

Figure 3D: V kappa 4 (Vx4) gene sequence

দ্র	GA CT		CA GT	Д
Q S P D S L A V S L G E BanII	GCCTGGGCGA CGGACCCGCT	N C R S S Q S V L Y S S PstI	TATAGCAGCA	O P
ᆸ	CTG	Ω.	TAG ATC	Q
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6 <i>1</i>	rga act	ı	GAGCGTGCTG	L A W Y Q Q K P G KpnI SexAI
	1997 1777	>	TGC	ъ Se
K	CTGGCGGTGA GACCGCCACT	Ø	GAGCGTGCTG CTCGCACGAC	×
H	CT		GA	Oi.
ß	AGC FCG	Q	ATTAACTGCA GAAGCAGCCA TAATTGACGT CTTCGTCGGT	O.
Ω	ATZ TA	Ω.	CAG( GTC(	) It
Δ	GATATCGTGA TGACCCAGAG CCCGGATAGC CTATAGCACT ACTGGGTCTC GGGCCTATCG	W.	GAAGCAGCCA	V X
S BanII	GAG CCC	R.	~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~	>
S Ba	3AG CTC	C ] PstI	CTGCA G	4
Q	CCA	<u>~</u> ₽	ACT IGA	H
<b>:</b>	TGACCCAGAG ACTGGGTCTC	Н	ATTAACTGCA TAATTGACGT	<b>&gt;</b> ⊣
Σ	T TO			<b>5</b>
V	~~~~~~ GATATCGTGA CTATAGCACT	E	ACGTGCGACC TGCACGCTGG	N N K N
	~~ ICG AGC	R A T	9090	<b>×</b>
D I EcoRV	~~~~~~ GATATC CTATAG	CK.	CGT	Z
μн	. 6 5		Ą	Z

AGGGCCTAGC 召 TCCCGGATCG AGAAACCAGG TCAGCCGCCG AGTCGGCGGC Ω Д ~~~~~~~ SanDI CTTTCGCCCC TCTTTGGTCC GAAAGCGGGG O Ŋ 口 ATCCACCCGT TAGGTGGGCA TGGTACCAGC ACCATGGTCG 召 Н Ŋ TTTATTGGGC CTATCTGGCG GATAGACCGC K 3 H TTTGATAATT ACAACAAAA AAACTATTAA TGTTGTTTT AseI 口 Ц X

AAATAACCCG

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Figure 3D: V kappa 4 (VK4) gene sequence (continued)

| G T D F T L T I S S | GCACTGATTT TACCCTGACC ATTTCGTCCC<br>CGTGACTAAA ATGGGACTGG TAAAGCAGGG | V Y Y С Q Q Н Y Т Т | GTGTATTATT GCCAGCAGCA TTATACCACC<br>CACATAATAA CGGTCGTCGT AATATGGTGG | T K V E I K R T BsiWI ~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~       |
|---------------------|----------------------------------------------------------------------|---------------------|----------------------------------------------------------------------|------------------------------------------------------------------|
| H                   | GCACT                                                                | X >                 | GTGTA                                                                | TACGAAAGT                                                        |
| F S G S G S G BamHI | TTTTAGCGGC TCTGGATCCG                                                | D V A               | BbsI<br>TGCAAGCTGA AGACGTGGCG<br>ACGTTCGACT TCTGCACCGC               | P P T F G Q G  MscI  CCGCCGACCT TTGGCCAGGG GGCGGCTGGA AACCGGTCCC |

|                                                                                | •                                  |            |                          | •           |                                 |               |                                       |
|--------------------------------------------------------------------------------|------------------------------------|------------|--------------------------|-------------|---------------------------------|---------------|---------------------------------------|
| <b>K</b>                                                                       | <u>ი</u> ი                         | <b>.</b> . | AGCAACTATG<br>TCGTTGATAC | <b>&gt;</b> | GCTGATTTAT<br>CGACTAAATA        | Н             | ì                                     |
| Q                                                                              | CAG                                | ≯          | CTZ                      | н           | TT'<br>PAA                      | G S<br>BamHI  | \<br>\<br>\<br>\                      |
| ڻ                                                                              | caggrcagcg<br>grccagrcgc           | Z          | CAA                      | J.          | TG2<br>AC1                      | ന പ്ല         | ì                                     |
| P<br>SexAI                                                                     | AC CAGGT<br>TG GTCCA               | Ŋ          | AG                       | , -         | S<br>S<br>S                     | ß             |                                       |
|                                                                                | ~ ∪ ⊕<br>H ⊕                       | ტ          | ည<br>ပို့                | H           | CT                              |               |                                       |
| Ø                                                                              | 200                                | Н          | CAACATTGGC<br>GTTGTAACCG | ×           | CGCCGAAACT<br>GCGGCTTTGA        | Ţ.            |                                       |
| Q                                                                              | 000                                |            | CA                       | Д           | 300<br>300<br>300               | CC.           |                                       |
| ഗ                                                                              | AGTGGCGCAC<br>TCACCGCGTG           | Z          | CAACATTGGC<br>GTTGTAACCG | A<br>BbeI   |                                 | Ω             |                                       |
| 1                                                                              |                                    | Ø          | AG                       |             | ( 00 )<br>( 00 )                | വ             |                                       |
| >                                                                              | GCCTTCAGTG<br>CGGAAGTCAC<br>Eco57I | Ŋ          | GCAGCAGCAG               | E           | CCCGGGACGG                      | >             |                                       |
| ß                                                                              | CTTCAG<br>GAAGTC<br>Eco57I         | ω ,        | AGC.                     | P G<br>XmaI | <br>                            | O             | 1                                     |
| വ                                                                              | 000<br>000<br>000<br>1             |            | GC7<br>CG1               | 면 X         | ~<br>000<br>000                 |               | \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ |
| Д                                                                              |                                    | ტ          | ტ<br>ე                   | Ъ           | TG                              | P S<br>Bsu36I | ;<br>;                                |
| ال ل                                                                           | AGC                                | W          | rag<br>atc               | O.          | AGT<br>TCA                      | 다<br>B        | }                                     |
| T.                                                                             | TGACCCAGCC<br>ACTGGGTCGG           | CSI        | TCGTGTAGCG<br>AGCACATCGC | Q           | .~~<br>CCAGCAGTTG<br>GGTCGTCAAC | 民             |                                       |
| gene si                                                                        | TGA<br>ACT                         | S<br>BssS  | TCGT                     |             | 200<br>000                      | Q             |                                       |
| (M)                                                                            | _                                  | Н          | ıc<br>AG                 | Y<br>KpnI   | GGTA<br>CCAT                    |               |                                       |
| bda 1<br>V                                                                     | GTC                                |            | CA,                      | Z<br>K      | rgg<br>ACC                      | Z             |                                       |
| V lam<br>S                                                                     | AGC                                | E          | GAC                      | Ŋ           | TGAGCTGGTA<br>ACTCGACCAT        | Z             |                                       |
| Figure 4A: V lambda 1 (V $\lambda$ 1) gene sequence Q S V $ m L$ $ m T$ $ m C$ | CAGAGCGTGC<br>GTCTCGCACG           | >          | TGTGACCATC<br>ACACTGGTAG | >           | TG?<br>AC1                      | Ω.            |                                       |
| Fig.                                                                           | . 00                               |            | •                        |             |                                 | ٠             |                                       |

Figure 4A: V lambda 1 (VA.1) gene sequence (continued)

| GCGGATCCAA<br>CGCCTAGGTT | S E D<br>BbsI | AGCGAAGACG                                     | V F G<br>TGTGTTTGGC<br>ACACAAACCG                           |                    |                          |
|--------------------------|---------------|------------------------------------------------|-------------------------------------------------------------|--------------------|--------------------------|
| GATCGTTTTA<br>CTAGCAAAAT | о<br>П        | GGGCCTGCAA<br>CCCGGACGTT                       | Q H Y T T P P P CAGCATTATA CCACCCGCC GTCGTAATAT GGTGGGGGGGG |                    |                          |
| AGGCGTGCCG               | T T           | TTGCGATTAC<br>AACGCTAATG                       | Q H Y T<br>CAGCATTATA (<br>GTCGTAATAT (                     | L G<br>MscI<br>~~~ | TCTTGGC<br>AGAACCG       |
| AGCGTCCCTC<br>TCGCAGGGAG | S<br>A<br>S   | AGCGCGAGCC                                     | Y C Q<br>TTATTGCCAG<br>AATAACGGTC                           | L T V<br>HpaI      | AGTTAACCGT<br>TCAATTGGCA |
| GATAACAACC<br>CTATTGTTGG | S<br>D        | AAGCGGCACC AGCGCGAGCC<br>TTCGCCGTGG TCGCGCTCGG | E A D Y<br>AAGCGGATTA<br>TTCGCCTAAT                         | G<br>T<br>K        | GGCGGCACGA               |

|                                      | •            |                                    |            |                                                |                                       |                                                |               |                          |
|--------------------------------------|--------------|------------------------------------|------------|------------------------------------------------|---------------------------------------|------------------------------------------------|---------------|--------------------------|
|                                      | Ø            | AG                                 |            | CT                                             | Н                                     | TT                                             | SHI           | TC<br>AG                 |
|                                      | Q            | CAG                                | Z          | raa<br>Att                                     | $oldsymbol{oldsymbol{arphi}}$         | rga<br>act                                     | G S<br>BamHI  | GGATC<br>CCTAG           |
|                                      | ٠<br>٢       | CAC                                | ×          | TA                                             | ᆡ                                     | GAC                                            | ω<br>L        |                          |
|                                      | P G<br>SexAI | CAGGTCAGAG<br>GTCCAGTCTC           | හ          | GGCTATAACT<br>CCGATATTGA                       | H                                     | ACTGATGATT<br>TGACTACTAA                       |               | TTAGCGGATC<br>AATCGCCTAG |
|                                      | Se<br>Se     | 1                                  | z h        |                                                | ×                                     |                                                | ĮΉ            |                          |
|                                      | Ω,           | TC?<br>AG1                         | Ŋ          | 999                                            | വ                                     | CGA                                            | 民             | CGT                      |
|                                      | Ŋ            | AGCGGCTCAC                         | >          | CGATGTGGGC<br>GCTACACCCG                       | A P<br>BbeI                           | AGGCGCCGAA<br>TCCGCGGCTT                       | Z             | AGCAACCGTT<br>TCGTTGGCAA |
|                                      | დ<br>დ       | )<br>(CG(                          | Ω          | GAC                                            | A E                                   | 000                                            | ഗ             | GC7                      |
|                                      | ,            | . –                                | Ŋ          |                                                | ×                                     |                                                |               |                          |
|                                      | >            | GTGTCAC                            |            | CAC<br>GTC                                     |                                       | GG2                                            | >             | GTC                      |
|                                      | တ            | CTTCAG<br>GAAGTC<br>Eco57I         | Q          | TAG<br>ATC                                     | P G<br>XmaI                           | 0<br>0<br>0<br>0<br>0<br>0                     |               | ~<br>380<br>380          |
|                                      | <b>⋖</b>     | AGCTTCAGTG<br>TCGAAGTCAC<br>Eco57I | H          | GTACTAGCAG<br>CATGATCGTC                       | •                                     | CATCCCGGGA<br>GTAGGGCCCT                       | S<br>191      | cercageceres cages       |
|                                      |              | A<br>T                             | O          | :                                              | 田                                     | CA                                             | P S<br>Bsu36I | CT<br>GA                 |
|                                      | Д            | 0<br>0<br>0<br>0                   | E          | 0<br>0<br>0<br>0                               | O1                                    | AG                                             | <u>д</u>      | ეეე<br> <br>  ტე         |
| nence                                | Q            | CAC                                |            | TAC                                            | α (                                   | CAGCAG<br>GTCGTC                               | 民             | CGI                      |
| sedne                                | H            | TGACCCAGCC<br>ACTGGGTCGG           | SSH<br>~~~ | TCGTGTACGG                                     | · · · · · · · · · · · · · · · · · · · | ACC<br>IGG                                     | Z             | GCAACCGTCC<br>CGTTGGCAGG |
| gene (                               | ,            | TG.                                | BSSS       | TC(<br>AG(                                     | W Y KpnI                              | GTAC<br>CATG                                   |               | 725                      |
| 2 (V)2                               | ij           | AC<br>TG                           | <b>⊢</b>   | TC<br>AG                                       | 3                                     | TG<br>AC                                       | W             | GA                       |
| nbda 🤅                               | A            | 0 0                                | E          | CCA                                            | Ø                                     | 16C                                            | D Q           | GT(                      |
| :Vlar                                | Ŋ            | AGC                                |            | TAC                                            | >                                     | TG2<br>AC1                                     | Q             | GA1<br>CT2               |
| Figure 4B: V lambda 2 (Vλ2) gene seq | Q            | CAGAGCGCAC<br>GTCTCGCGTG           | H          | CATTACCATC TCGTGTACGG<br>GTAATGGTAG AGCACATGCC | <b>≯</b>                              | ATGTGAGCTG GTACCAGCAG<br>TACACTCGAC CATGGTCGTC | $\succ$       | TATGATGTGA<br>ATACTACACT |
| Fig                                  |              |                                    |            |                                                | <b>r</b> -                            | 7 6                                            |               | L · 1~4                  |

Figure 4B: V lambda 2 (VA2) gene sequence (continued)

| SI  | AAG      | J.I.C                                  | ĮΤί                                    | דדי:<br>AA                                                                                            |                                                                                                                                                                                      |                                                                                                                                                                                                                |
|-----|----------|----------------------------------------|----------------------------------------|-------------------------------------------------------------------------------------------------------|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| 西   | 3G.Z     |                                        | 5                                      | rg1<br>AC2                                                                                            |                                                                                                                                                                                      |                                                                                                                                                                                                                |
|     | CC       | CG                                     |                                        | TG.                                                                                                   |                                                                                                                                                                                      |                                                                                                                                                                                                                |
|     | CAA      | GTT                                    |                                        | 0<br>0<br>0<br>0<br>0                                                                                 |                                                                                                                                                                                      | ٠                                                                                                                                                                                                              |
|     | TG       | ;AC                                    | വ                                      | ည<br>ညီ ဗ္                                                                                            |                                                                                                                                                                                      |                                                                                                                                                                                                                |
|     | 300      | C<br>C<br>C<br>C                       | E                                      | ACC<br>IGG                                                                                            |                                                                                                                                                                                      |                                                                                                                                                                                                                |
|     | CGC      | U<br>U                                 | £                                      | 200                                                                                                   |                                                                                                                                                                                      |                                                                                                                                                                                                                |
|     | 'AG      | D'L                                    |                                        | TA                                                                                                    |                                                                                                                                                                                      |                                                                                                                                                                                                                |
|     |          |                                        |                                        |                                                                                                       |                                                                                                                                                                                      |                                                                                                                                                                                                                |
|     | CAT      | ZT.                                    | <del>     </del>                       | ATT<br>PAP                                                                                            | Ω<br>Ω<br>I                                                                                                                                                                          | ~~~~<br>TGGC<br>ACCG                                                                                                                                                                                           |
|     | ACC      | JG(                                    | +44                                    | 200<br>200                                                                                            | i<br>Z                                                                                                                                                                               | TTC<br>AAC                                                                                                                                                                                                     |
|     | TG.      | AC                                     | Q                                      | CA(<br>GT(                                                                                            | • •                                                                                                                                                                                  | TC.<br>AG.                                                                                                                                                                                                     |
|     | SCC      | GG.                                    | Q                                      | AG<br>TC                                                                                              | >                                                                                                                                                                                    | CGTTCTTGGC<br>GCAAGAACCG                                                                                                                                                                                       |
|     |          |                                        |                                        |                                                                                                       | உ                                                                                                                                                                                    |                                                                                                                                                                                                                |
|     | CGZ      | 3C1                                    |                                        | IGC<br>ACC                                                                                            | a<br>H                                                                                                                                                                               | CGAAGTTAAC<br>GCTTCAATTG                                                                                                                                                                                       |
|     | SS       | Ü<br>Ü                                 | ≻                                      | AT                                                                                                    | L<br>HD                                                                                                                                                                              | TT.<br>AA'                                                                                                                                                                                                     |
|     | :AC      | TG                                     | <b>k</b> .                             | TT                                                                                                    | ×                                                                                                                                                                                    | AAG                                                                                                                                                                                                            |
|     | AAC      | rTG                                    | ×                                      | rt?<br>AAT                                                                                            |                                                                                                                                                                                      | 767<br>301                                                                                                                                                                                                     |
|     | 2        |                                        | D                                      |                                                                                                       | E                                                                                                                                                                                    |                                                                                                                                                                                                                |
|     | GGC      | CC<br>CC                               |                                        | GG2                                                                                                   |                                                                                                                                                                                      | 250                                                                                                                                                                                                            |
|     | Ď,       | C<br>G                                 | Ø                                      | ည္သ                                                                                                   | - <b>h</b>                                                                                                                                                                           | 900                                                                                                                                                                                                            |
| HI  | AA       | LL                                     | 田出                                     | AAS<br>TT                                                                                             | U                                                                                                                                                                                    | 900                                                                                                                                                                                                            |
| Ban | ~<br>CAA | GTJ                                    | D<br>Bbs                               | ACG<br>TGG                                                                                            | Ŋ                                                                                                                                                                                    | GGCGGCGGCA                                                                                                                                                                                                     |
|     | BamHI    | GCGGC AACACCGCGA GCCTGACCAT TAGCGGCCTG | GCGGC AACACCGCGA GCCTGACCAT TAGCGGCCTG | GCGGC AACACCGCGA GCCTGACCAT TAGCGGCCTG CGCCG TTGTGGCGCT CGGACTGGTA ATCGCCGGAC A D Y Y C Q Q H Y T T P | GCGGC AACACCGCGA GCCTGACCAT TAGCGGCCTG CGCCG TTGTGGCGCCT CGGACTGGTA ATCGCCGGAC A D Y Y C Q Q H Y T T P GCGGA TTATTATTGC CAGCAGCATT ATACCACCCC CGCCT AATAATAACG GTCGTCGTAA TATGGTGGGG | GCGGC AACACCGCGA GCCTGACCAT TAGCGGCCTG CGCCCG TTGTGGCGCT CGGACTGGTA ATCGCCGGAC A D Y Y C Q Q H Y T T P GCGGA TTATTATTGC CAGCAGCATT ATACCACCCC CGCCT AATAATAACG GTCGTCGTAA TATGGTGGGG G T K L T V L G HDAI MSCI |

Figure 4C: V lambda 3 (VA3) gene sequence

| H                      | AC                                 |              | CT                       | Д                     | AT.                      |
|------------------------|------------------------------------|--------------|--------------------------|-----------------------|--------------------------|
| Q                      | CAG                                | ÓΊ           | GAG                      | Ω                     | ATG                      |
| σ                      | ~ Z<br>G T<br>C A                  | 4            | )<br>(0<br>(0<br>(0      |                       | TG                       |
| S V A P G Q T<br>SexAI | AC CAGGTCAGAC<br>IG GTCCAGTCTG     | G D K Y A S  | TACGCGAGCT<br>ATGCGCTCGA | Y D D                 | TTATGATGAT<br>AATACTACTA |
| გ                      | }                                  | ₩.           | A.T.                     | н                     |                          |
| 4                      | , GC2<br>, CG3                     | 택            | TAZ<br>ATJ               | >                     | TG2<br>AC1               |
| >                      | GTT                                | Д            | CGA                      | . ٦                   | rgg<br>ACC               |
| W                      | AGCGTTGCAC<br>TCGCAACGTG           | O            | GGGCGATAAA<br>CCCGCTATTT | <b>—</b>              | TTCTGGTGAT<br>AAGACCACTA |
|                        |                                    | L)           |                          | Q A P V L V I<br>Bbei |                          |
| P S V                  | AGT<br>CAC                         |              | S<br>S<br>S<br>S<br>S    | Д.,                   | CA(                      |
| Ŋ                      | CTTCAG<br>GAAGTC<br>Eco57I         | <b>A</b>     | TGC                      | A<br>BbeI             | 992922<br>229295         |
|                        | GCCTTCAGTG<br>CGGAAGTCAC<br>Eco57I | D A L        | GCGATGCGCT<br>CGCTACGCGA | ) M                   | CAGGCGCCAG               |
| а<br>О                 |                                    | <sub>ල</sub> |                          |                       |                          |
| щ                      | TGACCCAGCC<br>ACTGGGTCGG           | ω<br>O       | TCGTGTAGCG               | P G<br>XmaI           | GAAACCCGGG               |
| Q                      | CAC                                |              | TAC                      | P G<br>XmaI           | 000                      |
| H                      | ACC                                | S C<br>BssSI | GTG                      |                       | AAC<br>ITG               |
| ᆸ                      | A P C                              | SCBSSSI      | TC A                     | ×                     | GA                       |
| •                      | AC<br>TG                           | H            |                          | о<br>о                | CA                       |
| 团                      | rga<br>act                         | <b>K</b>     | STA<br>CAT               | Q,                    | CAG                      |
| <b>&gt;</b>            | TAT                                |              | CGCGCGTATC               | W Y<br>KpnI           | ACC<br>TGC               |
| Ŋ                      | AGCTATGAAC<br>TCGATACTTG           | <b>A</b>     | CGCGCGTATC               | W<br>Kp               | GGTACCAGCA               |
|                        | • –                                |              | -                        | · ·                   | - <b>-</b>               |

Figure 4C: V lambda 3 (VA.3) gene sequence (continued)

G CCAACAGCGG GGTTGTCGCC K S 田 Z Д ~~~~~~ BbsI BamHI TTTAGCGGAT AAATCGCCTA 口 Ö K S Ø ſщ GGGCCTTGCG CCCGGAACGC H 召 <u>ෆ</u> 团 Ø Д Н GGAGTCCGTA CCTCAGGCAT Н C ~~~~~~ Bsu36I П Ŋ 드 Д AGACTGGCAG TCTGACCGTC Ø K H Д Z Ŋ

GACGAAGCGG CTGCTTCGCC TGGCGGCGGC ACCGCCGCCG ტ O TCAGGCGGAA AGTCCGCCTT CGCCTGTGTT GCGGACACAA Д Д ATATGGTGGG TTAGCGGCAC AATCGCCGTG TATACCACCC H E ACCCTGACCA Q Q H CCAGCAGCAT TGGGACTGGT GGTCGTCGTA CAACACCGCG GTTGTGGCGC TAATAATAAC ပ ATTATTATTG

T K L T V L G
HpaI MscI

ACGAAGTTAA CCGTTCTTGG C TGCTTCAATT GGCAAGAACC G

|                                      | Q           | CAG                                                                        | 4            | AGCTATGCGA<br>TCGATACGCT | Ö           | GATGGGCGGC                                                                                  | TTCAGGGCCG                                | 긔             |
|--------------------------------------|-------------|----------------------------------------------------------------------------|--------------|--------------------------|-------------|---------------------------------------------------------------------------------------------|-------------------------------------------|---------------|
|                                      | Ø           | CGGGCAGCAG<br>GCCCGTCGTC                                                   | Ą            | TGC                      | ტ           | 990                                                                                         | ပ ပို ပို                                 |               |
|                                      | O           | 00<br>00<br>00<br>00<br>00<br>00<br>00<br>00<br>00<br>00<br>00<br>00<br>00 | ×            | CTA<br>SAT               | ×           | rgg<br>ACC                                                                                  | O<br>CAG<br>GTC                           | 闰             |
|                                      |             | 200                                                                        | വ            | AGC                      | 4           | GA!<br>CT2                                                                                  | TT(<br>AA(                                | Σ             |
|                                      | Д           | AC<br>FG                                                                   | W            | ე<br>ე                   | 3           | IG<br>AC                                                                                    | GH C                                      | <b>&gt;</b>   |
|                                      | ×           | AA/<br>TT:                                                                 |              | TA(<br>AT(               | 田上          | ZAG.                                                                                        | K<br>AA(<br>TT(                           |               |
|                                      | ×           | AAA<br>I'T'I                                                               | Ţ'n          | rtt<br>AAA               | L E<br>XhoI | CTCGA                                                                                       | A Q<br>CGCAG<br>GCGTC                     | K             |
|                                      | >           | GTGAAAAAAC<br>CACTTTTTTG                                                   | E            | CACTTTTAGC<br>GTGAAAATCG |             | GTCTCGAGTG<br>CAGAGCTCAC                                                                    | A'QK<br>GCGCAGAAGT<br>CGCGTCTTCA          | H             |
|                                      |             |                                                                            | Check        |                          | Ö           | _                                                                                           |                                           | Ŋ             |
| ٠                                    | 时           | GA                                                                         |              | AGG                      | Q           | AG                                                                                          | Y<br>YTA:                                 |               |
|                                      | Æ           | 999                                                                        | S G<br>BSPEI | 000                      | r<br>U      | 000<br>0000                                                                                 | N<br>AAC<br>TTC                           | H             |
|                                      | O           | TGGCGCGGAA<br>ACCGCGCCTT                                                   | ഗ <u>എ</u> ≀ |                          | Q           | CCAAGCC CCTGGGCAGG                                                                          | A N Y<br>GGCGAACTAC<br>CCGCTTGATG         | Q             |
|                                      |             | •                                                                          | 4            |                          | XH          | ₹                                                                                           | _                                         | 田             |
| nce                                  | W           | TGGTTCAGTC<br>ACCAAGTCAG                                                   | ×            | AGCTGCAAAG<br>TCGACGTTTC | A<br>BstXI  | CCAAGCC                                                                                     | T<br>CAC<br>GTG                           | Д             |
| ene sequence                         | Q           | CA(                                                                        |              | CAZ                      | OI .        | AA(                                                                                         | 0<br>0<br>0<br>0<br>0<br>0<br>0<br>0<br>0 |               |
| gene                                 | >           | GTT                                                                        | Ö            | CTG                      | <b>K</b>    | 3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3 | F<br>TTT<br>AAA                           | A             |
| VH1A)                                | 2 3         | TGGTTCAGTC<br>ACCAAGTCAG                                                   | <b>ග</b>     | AG                       |             | GCGCCAAGCC                                                                                  | I F G T<br>TTTTGGCAC<br>AAAAACCGTG        | H             |
| n 1A (                               | Q L<br>MfeI | AT<br>T.A                                                                  | >            | TG<br>AC                 | >           | GT                                                                                          |                                           | н             |
| ,<br>γ chai                          | Ø Z ∑       | SCA                                                                        | ¥            | AAG<br>FTC               | S           | TTAGCTGGGT<br>AATCGACCCA                                                                    | PCC.                                      |               |
| / heav                               | >           | GTC                                                                        |              | GAZ                      | ഗ           | , CG.                                                                                       | I<br>TAT                                  | Щ             |
| Figure 5A: V heavy chain 1A (VH1A) g | Q           | CAGGTGCAAT<br>GTCCACGTTA                                                   | >            | CGTGAAAGTG<br>GCACTTTCAC | H           | TTAGCTGGGT<br>AATCGACCCA                                                                    | I I P<br>ATTATTCCGA<br>TAATAAGGCT         | V T<br>BstEII |
| Figu                                 |             | . <del>-</del>                                                             |              |                          |             |                                                                                             |                                           |               |
|                                      |             |                                                                            |              |                          |             |                                                                                             |                                           |               |

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| :hain 1,A (VH1A) gene sequence (continued) |
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| Figure                                     |
| :-                                         |
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| GGTGACCATT ACCGCGGATG AAAGCACCAG CACCGCGTAT ATGGAACTGA<br>CCACTGGTAA TGGCGCCTAC TTTCGTGGTC GTGGCGCATA TACCTTGACT | S L R S E D T A V Y Y C A R W G Eagl Eagl | GCAGCCTGCG TAGCGAAGAT ACGGCCGTGT ATTATTGCGC GCGTTGGGGC<br>CGTCGGACGC ATCGCTTCTA TGCCGGCACA TAATAACGCG CGCAACCCCG | DGFYAMDYWGQGTLVT | GGCGATGGCT TTTATGCGAT GGATTATTGG GGCCAAGGCA CCCTGGTGAC<br>CCGCTACCGA AAATACGCTA CCTAATAACC CCGGTTCCGT GGGACCACTG | V S S<br>BlpI | GGTTAGCTCA G |
|------------------------------------------------------------------------------------------------------------------|-------------------------------------------|------------------------------------------------------------------------------------------------------------------|------------------|------------------------------------------------------------------------------------------------------------------|---------------|--------------|
| GGTGACC                                                                                                          |                                           | GCAGCC1<br>CGTCGGA                                                                                               | D<br>D           | GGCGATC                                                                                                          | ><br>>        | GGTTAG       |

Figure 58: V heavy chain 18 (VH1B) gene sequence

| w                                     | AG                         |              | TA<br>AT                 | M              | gg<br>CC                                       | 3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3 |
|---------------------------------------|----------------------------|--------------|--------------------------|----------------|------------------------------------------------|---------------------------------------------------------------------------------------------|
| <b>₹</b>                              | CGGGCGCGAG                 | X<br>X       | AGCTATTATA<br>TCGATAATAT | M<br>G<br>M    | GATGGGCTGG<br>CTACCCGACC                       | F Q G R<br>TTCAGGGCCG<br>AAGTCCCGGC                                                         |
| <u>U</u>                              | 0000                       | ≯            | TAZ                      | _              | 999                                            | D D D D D D D D D D D D D D D D D D D                                                       |
|                                       | 000                        | S            | AGC<br>ICG               | Σ              | GAT                                            | TTC                                                                                         |
| Д                                     |                            | <b>r</b> .   | •                        | Z              |                                                |                                                                                             |
| ×                                     | AAA<br>I'T'I               | Н            | rac<br>Atg               | ы<br>ы н а     | AGT<br>ICA                                     | K<br>AAG<br>TTC                                                                             |
| <b>×</b>                              | AA                         | T<br>F       | TT                       | L E<br>XhoI    | S S                                            | O S T                                                                                       |
| V K K P                               | GTGAAAAAAC<br>CACTTTTTTG   | Ħ            | TACCTTTACC<br>ATGGAAATGG | HX             | GTCTCGAGTG                                     | A Q K F<br>GCGCAGAAGT<br>CGCGTCTTCA                                                         |
| •                                     | •                          |              | •                        | Ö              |                                                |                                                                                             |
| 时                                     | CGGCGCGGAA<br>GCCGCGCCTT   | ≯<br>⊢ ≀     | CCTCCGGATA<br>GGAGGCCTAT | р G D          | CCTGGGCAGG                                     | T N Y<br>CACGAACTAC<br>GTGCTTGATG                                                           |
| ව<br>අ                                | ,<br>,<br>,<br>,<br>,<br>, | S G<br>BSPEI | 3000                     | · .            | 2002                                           | N<br>NAC<br>PTG                                                                             |
| <b>U</b>                              | 000                        | လည္          | TCC                      |                | TGG                                            | T<br>CGA                                                                                    |
|                                       | 0<br>0<br>0                |              | 0 0                      | KT P           |                                                | CA                                                                                          |
| Ø                                     | AG<br>TC                   | N<br>M       | AG                       | A P G<br>BstxI | S<br>S<br>S<br>S<br>S                          | 0<br>0<br>0<br>0<br>0                                                                       |
| S<br>O                                | TGGTTCAGAG<br>ACCAAGTCTC   | X            | AGCTGCAAAG<br>TCGACGTTTC | O A Bst        |                                                | S G G<br>ATAGCGGCGG<br>TATCGCCGCC                                                           |
| >                                     | TTC                        | Ö            | TGC                      |                | CC7                                            | န<br>ငြင်င                                                                                  |
| l<br>l                                | TGGT<br>ACCA               | ഗ            | AGC<br>ICG               | K.             | )<br>9<br>9<br>9<br>9<br>9<br>9<br>9<br>9<br>9 | ATA<br>TAT                                                                                  |
| ДН≀                                   |                            |              |                          | >              |                                                | Z                                                                                           |
| M M M M M M M M M M M M M M M M M M M | CAA                        | <b>&gt;</b>  | AGT                      | M              | 0000                                           | P<br>CCC<br>GGC                                                                             |
| >                                     | TG<br>AC                   | ×            | SAA                      | Ħ              | ACT<br>FGA                                     | N<br>AAC<br>FTG                                                                             |
| Ol                                    | CAGGTGCAAT<br>GTCCACGTTA   | >            | CGTGAAAGTG<br>GCACTTTCAC |                | TGCACTGGGT<br>ACGTGACCCA                       | I N P<br>ATTAACCCGA<br>TAATTGGGCT                                                           |
|                                       | O O                        |              | O O                      | Σ              | HK                                             | AH                                                                                          |
|                                       |                            |              |                          |                |                                                |                                                                                             |

|                                    | ᆸ             | TGA<br>ACT               | to             | 522<br>255                                                                      | <b>[-1</b>         | GAC                          |
|------------------------------------|---------------|--------------------------|----------------|---------------------------------------------------------------------------------|--------------------|------------------------------|
|                                    | ED<br>L       | AAC<br>ITG               | R<br>R         | IGG<br>ACC                                                                      | >                  | SGT                          |
|                                    | ы<br>Ж        | ATGGAACTGA<br>TACCTTGACT | <b>~</b> , ,   | GCGTTGGGGC                                                                      | G Q G T L V T Styl | CCCTGGTGAC                   |
|                                    | •             |                          | H H H          |                                                                                 | E                  |                              |
|                                    | ×             | CACCGCGTAT               | C A<br>BSSHII  | ATTATTGCGC<br>TAATAACGCG                                                        | ַ ט                | GGCCAAGGCA<br>CCGGTTCCGT     |
|                                    | T A Y         | 999                      |                | TTG                                                                             | O G<br>Styl        | AAG<br>TTC                   |
|                                    | H             | ACC<br>IGG               | X X >          | LTA<br>AAT                                                                      | (n) {              | 1000<br>1000<br>1000<br>1000 |
|                                    |               |                          | ≯              |                                                                                 |                    |                              |
|                                    | ß             | TAG<br>ATC               | >              | TGT                                                                             | D Y W              | TGG<br>ACC                   |
|                                    | Н             | CAT                      | Bagi<br>~~~~~~ | 30<br>0<br>0<br>0                                                               | <b>≯</b>           | rat<br>Ata                   |
| red)                               | Ø             | CCAGCATTAG<br>GGTCGTAATC |                | ACGGCCGTGT<br>TGCCGGCACA                                                        | Ω                  | GGATTATTGG<br>CCTAATAACC     |
| gene sequence (continued)          | EH            |                          | [-  ≀          | •                                                                               |                    |                              |
| ence (c                            | 0             | ATA<br>FAT               | Ω              | SAT                                                                             | Σ                  | SAT                          |
| e segu                             | A<br>U        | TG                       | EI<br>CI       | AA(<br>TT(                                                                      | Y A M              | )<br>)<br>)<br>)<br>)        |
| B) gen                             |               | ACCCGTGATA<br>TGGGCACTAT | Ω.             | TAGCGAAGAT<br>ATCGCTTCTA                                                        | ×                  | TTTATGCGAT<br>AAATACGCTA     |
| 3 (VH1                             | H             | AC                       |                | TA                                                                              | Гц                 | TT                           |
| Jain 1E                            | Σ             | TG                       | K<br>K         | 30<br>30<br>30<br>30<br>30<br>30<br>30<br>30<br>30<br>30<br>30<br>30<br>30<br>3 |                    | SCT                          |
| eavy cł                            | ĦН            | GGTGACCATG<br>CCACTGGTAC | Н              | GCAGCCTGCG<br>CGTCGGACGC                                                        | <sub>C</sub>       | GGCGATGGCT<br>CCGCTACCGA     |
| B: V h                             | V T<br>BstEII | GGTGACC                  | w              | AGC<br>TCG                                                                      | Q                  | CGA                          |
| Figure 5B: V heavy chain 1B (VH1B) | BS            | 000<br>000               | တ              | G<br>G                                                                          | Ŋ                  | 0<br>0<br>0                  |
| ΙÏ                                 |               |                          |                |                                                                                 |                    |                              |

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V S S BlpI CCAATCGAGT C

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TATAGCACCA GCCTGAAAAC ATATCGTGGT CGGACTTTTG

TGATAAGTAT ACTATTCATA

ATTGGGATGA TAACCCTACT

GCTCTGATTG

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| w                 | AG                                                                                          |             | GA                       | Æ          | ດ<br>ດ                         | K 0 0                                                                                       |
|-------------------|---------------------------------------------------------------------------------------------|-------------|--------------------------|------------|--------------------------------|---------------------------------------------------------------------------------------------|
| <mark>ෆ</mark>    | CGGGCGGCAG<br>GCCCGCCGTC                                                                    | X A         | AGCTATGCGA<br>TCGATACGCT |            | GGTGAGCGCG<br>CCACTCGCGC       | K G R<br>TGAAAGGCCG<br>ACTTTCCGGC                                                           |
| <sub>.</sub><br>ບ | 000                                                                                         | Ħ           | TA                       | w          | GAC                            | K<br>AA(                                                                                    |
|                   | 000                                                                                         | W           | AGC<br>ICG               | >          | GGT                            | IGA                                                                                         |
| Д                 | •                                                                                           |             | -                        | 3          |                                | >                                                                                           |
| Q                 | CAA                                                                                         | ·Ø          | TAGATC                   | щH         | AGT<br>ICA                     | s<br>AGC<br>ICG                                                                             |
| L V Q P           | CTGGTGCAAC<br>GACCACGTTG                                                                    | E<br>E<br>S | TACCTTTAGC<br>ATGGAAATCG | L E        | GTCTCGAGTG                     | A D S<br>GCGGATAGCG<br>CGCCTATCGC                                                           |
| H                 | TGG                                                                                         | H           | ACC<br>TGG               |            | TCT                            | A<br>CGG<br>GCC                                                                             |
|                   |                                                                                             | দ           |                          | O          |                                | •                                                                                           |
| Ö                 | ອວວອວວອວວອ                                                                                  |             | CCTCCGGATT<br>GGAGGCCTAA | ×          | CCTGGGAAGG                     | T Y Y<br>CACCTATTAT<br>GTGGATAATA                                                           |
| <sub>ອ</sub><br>ອ | 360                                                                                         | S<br>BSPEI  | 900                      | _          | SGA                            | Y<br>FAT<br>ATA                                                                             |
| Ö                 | S<br>S<br>S<br>S<br>S<br>S<br>S<br>S<br>S<br>S<br>S<br>S<br>S<br>S<br>S<br>S<br>S<br>S<br>S | SA          | TC(                      | <b>O</b>   | T G                            | FCC.                                                                                        |
|                   | 000                                                                                         | Æ           | ပ<br>ပ                   | A X        | 000                            | CA                                                                                          |
| ß                 | AAG                                                                                         | A<br>A      | 0<br>0<br>0<br>0         | A<br>BstXI | CCAAGCC CCTGG<br>GGTTCGG GGACC | S<br>CAG                                                                                    |
| 国                 | TGGTGGAAAG<br>ACCACCTTTC                                                                    |             | AGCTGCGCGG<br>TCGACGCGCC | O H        | GCGCCAAGCC                     | GCGCGGCAG                                                                                   |
| >                 | STG                                                                                         | Ö           | CTG                      | <b>A</b>   | 305<br>206                     | 3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3<br>3 |
| 1                 |                                                                                             |             | AG(                      | <b>—</b>   | S<br>S<br>S<br>S<br>S          | , .<br>,                                                                                    |
| Q L<br>Mfei       | AT<br>TA                                                                                    | H           | TG<br>AC                 | >          | GT                             | S<br>TA<br>AT                                                                               |
| OZ 1              | CGT                                                                                         |             | GTC                      | M          | 766<br>ACC                     | 0<br>0<br>0<br>0<br>0<br>0<br>0                                                             |
| >                 | GTC                                                                                         | R.          | ညည                       | W          | GC.                            | S<br>'AG(                                                                                   |
| 臼                 | GAAGTGCAAT<br>CTTCACGTTA                                                                    | 니           | CCTGCGTCTG<br>GGACGCAGAC | Σ          | TGAGCTGGGT<br>ACTCGACCCA       | I S G<br>ATTAGCGGTA<br>TAATCGCCAT                                                           |
|                   |                                                                                             |             |                          | · -•       |                                | , -                                                                                         |

GGTTAGCTCA CCAATCGAGT

Figure 5D: V heavy chain 3 (VH3) gene sequence (continued)

| gene sequence |
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| Q   V   Q   L   Q   E   S   G   P   G   L   V   K   P   S   E   T                                                                                                                                                                                                                                                                                         |            |                  |                                             |                           |            |                                         |                |                 |
|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------------|------------------|---------------------------------------------|---------------------------|------------|-----------------------------------------|----------------|-----------------|
| TGCAAGAAAG TGGTCCGGGC CTGGTGAAAC ACGTTCTTTC ACCAGGCCCG GACCACTTTG ACGTCCTTTC ACCAGGCCCG GACCACTTTG  T C T V S G G S I S BSPEI ACCTGCACCG TTTCCGGAGG CAGCATTAGC TGGACGTGGC AAAGGCCTCC GTCGTAATCG AGCGTCGC CCTGGGAAGG GTCTCGAGTG AGCGGTCGGC GGACCCTTCC CAGAGCTCAC S G S T N Y N P S L K GCGCCAGCC CAACTATAAT GGCTCGGACT GCGCGCGCCGGGC GTTGATATTA GGCTCGGACT | H          | AC               |                                             | TT                        | ⋈          | AT<br>TA                                | > I            | ~~<br>GT<br>CA  |
| TGCAAGAAAG TGGTCCGGGC CTGGTGAAAC ACGTTCTTTC ACCAGGCCCG GACCACTTTG ACGTCCTTTC ACCAGGCCCG GACCACTTTG  T C T V S G G S I S BSPEI ACCTGCACCG TTTCCGGAGG CAGCATTAGC TGGACGTGGC AAAGGCCTCC GTCGTAATCG AGCGTCGC CCTGGGAAGG GTCTCGAGTG AGCGGTCGGC GGACCCTTCC CAGAGCTCAC S G S T N Y N P S L K GCGCCAGCC CAACTATAAT GGCTCGGACT GCGCGCGCCGGGC GTTGATATTA GGCTCGGACT | 臼          | GAA              |                                             | TTA                       | ტ          | GCT                                     | R<br>stE       | ~<br>000<br>000 |
| TGCAAGAAAG TGGTCCGGGC CTGGTGAAAC ACGTTCTTTC ACCAGGCCCG GACCACTTTG ACGTCCTTTC ACCAGGCCCG GACCACTTTG  T C T V S G G S I S BSPEI ACCTGCACCG TTTCCGGAGG CAGCATTAGC TGGACGTGGC AAAGGCCTCC GTCGTAATCG AGCGTCGC CCTGGGAAGG GTCTCGAGTG AGCGGTCGGC GGACCCTTCC CAGAGCTCAC S G S T N Y N P S L K GCGCCAGCC CAACTATAAT GGCTCGGACT GCGCGCGCCGGGC GTTGATATTA GGCTCGGACT | w          | AGC<br>ICG       | ×                                           | CTA                       |            | LTG                                     |                | 16C             |
| TGCAAGAAG TGGTCCGGGC CTGGTGAAAC ACGTTCTTTC ACCAGGCCCG GACCACTTTG T C T V S G G S I S BSPEI  ACCTGCACCG TTTCCGGAGG CAGCATTAGC TGGACGTGGC AAAGGCCTCC GTCGTAATCG R Q P G K G L E W BStXI XhoI  TCGCCAGCG CCTGGGAAGG GTCTCGAGTG AGCGTCGGC GACCCTTCC CAGAGCTCAC  S G S T N Y N P S L  GCGCAGCAC CAACTATAAT GGCTCGAGTG CGCCGTCGTG GTTGATATTA GGCTCGGACT         | 0.         | CG/2             | Ω.                                          | AGO                       | ••         | GA5<br>CT2                              |                | AAA<br>TTT      |
| TGCAAGAAG TGGTCCGGGC ACGTTCTTTC ACCAGGCCCG TT V S G G BSPEI ACCTGCACGG TTTCCGGAGG TGGACGTGGC AAAGGCCTCC ACCTGCACCG CTTGGAAGG AGCGCTCGC GGACCTTCC AGCGCTCGC GGACCTTCC AGCGCTCGG GGACCCTTCC AGCGCTCGT N Y N GCGCCAGCAC CAACTATAAT CGCCGCCGTCGTG GTTGATATTA                                                                                                  |            | PG<br>FG         | ro.                                         | $\mathcal{O} \mathcal{O}$ | 3          | PG<br>C                                 | <b>14</b>      | A T             |
| TGCAAGAAG TGGTCCGGGC ACGTTCTTTC ACCAGGCCCG TT V S G G BSPEI ACCTGCACGG TTTCCGGAGG TGGACGTGGC AAAGGCCTCC ACCTGCACCG CTTGGAAGG AGCGCTCGC GGACCTTCC AGCGCTCGC GGACCTTCC AGCGCTCGG GGACCCTTCC AGCGCTCGT N Y N GCGCCAGCAC CAACTATAAT CGCCGCCGTCGTG GTTGATATTA                                                                                                  | ×          | AAZ              |                                             | TA(<br>AT(                | 田王         | ~~<br>AG1<br>TC2                        | ᆸ              | CTC             |
| TGCAAGAAG TGGTCCGGGC ACGTTCTTTC ACCAGGCCCG TT V S G G BSPEI ACCTGCACGG TTTCCGGAGG TGGACGTGGC AAAGGCCTCC ACCTGCACCG CTTGGAAGG AGCGCTCGC GGACCTTCC AGCGCTCGC GGACCTTCC AGCGCTCGG GGACCCTTCC AGCGCTCGT N Y N GCGCCAGCAC CAACTATAAT CGCCGCCGTCGTG GTTGATATTA                                                                                                  | >          | GTG              | Н                                           | CAT                       | г<br>Хро   | ~~~<br>ICG<br>AGC                       | ഗ              | AGC<br>ICG      |
| TGCAAGAAG TGGTCCGGGC ACGTTCTTTC ACCAGGCCCG TT V S G G BSPEI ACCTGCACGG TTTCCGGAGG TGGACGTGGC AAAGGCCTCC ACCTGCACCG CTTGGAAGG AGCGCTCGC GGACCTTCC AGCGCTCGC GGACCTTCC AGCGCTCGG GGACCCTTCC AGCGCTCGT N Y N GCGCCAGCAC CAACTATAAT CGCCGCCGTCGTG GTTGATATTA                                                                                                  | ᆸ          | CTG              | Ŋ                                           | CAG                       |            | GAG                                     | Д              | 3007            |
| L Q E S TGCAAGAAAG ACGTTCTTTC TGGACGTGGC TGGACGTGGC ACCTGCAGCCG AGCGGTCGGC AGCGGTCGGC AGCGCTCGTGCCC AGCGCTCGTGCCCCCCCCCC                                                                                                                                                                                                                                  |            |                  | O                                           |                           | Ŋ          | •                                       |                |                 |
| L Q E S TGCAAGAAAG ACGTTCTTTC TGGACGTGGC TGGACGTGGC ACCTGCAGCCG AGCGGTCGGC AGCGGTCGGC AGCGCTCGTGCCC AGCGCTCGTGCCCCCCCCCC                                                                                                                                                                                                                                  | U          | 900              | H (                                         | AAG                       | ×          | AAG<br>TTC                              | Z              | AA!<br>TTI      |
| L Q E S TGCAAGAAAG ACGTTCTTTC TGGACGTGGC TGGACGTGGC ACCTGCAGCCG AGCGGTCGGC AGCGGTCGGC AGCGCTCGTGCCC AGCGCTCGTGCCCCCCCCCC                                                                                                                                                                                                                                  | <u>ρ</u> , | )<br>)<br>)<br>) | gds<br>×××××××××××××××××××××××××××××××××××× | 900                       | ប          | ~<br>GG <u>/</u>                        | ₩              | TAJ<br>AT       |
| L Q E S TGCAAGAAAG ACGTTCTTTC TGGACGTGGC TGGACGTGGC ACCTGCAGCCG AGCGGTCGGC AGCGGTCGGC AGCGCTCGTGCCC AGCGCTCGTGCCCCCCCCCC                                                                                                                                                                                                                                  | O          | GGT              |                                             | LTC                       | 0          | A C                                     | Z              | AAC             |
| L Q E TGCAAGA ACGTTCT TGGACGT R Q R Q TCGCCAG AGCGGTC S G S                                                                                                                                                                                                                                                                                               |            | T<br>A           | >                                           | T \                       | X          |                                         |                | C2<br>G1        |
| L Q E TGCAAGA ACGTTCT TGGACGT R Q R Q TCGCCAG AGCGGTC S G S                                                                                                                                                                                                                                                                                               | S          | AG               |                                             | )<br>(GC                  | P<br>SS t  | 000                                     | <del>[ '</del> | AC              |
| TGCAAAACGTTTAGACTTAGACTTGAACGGGGGGGGGGG                                                                                                                                                                                                                                                                                                                   | 回          | SAA              | H                                           | CAC                       | Д<br>С     | 4 6 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 | ß              | AGC<br>ICG      |
| i i i i i i i i i i i i i i i i i i i                                                                                                                                                                                                                                                                                                                     | Q          | AA(<br>TT(       | Ü                                           |                           |            |                                         | <b>U</b>       | 200             |
| i i i i i i i i i i i i i i i i i i i                                                                                                                                                                                                                                                                                                                     |            | IGC<br>ACG       | H                                           | ACC<br>TGG                | щ          | TCG<br>AGC                              |                | 300             |
| CAGGTGCAAGTCCACGGACCTCGCACGGACCCTGAGCCTGGACTCGGACCTGACCTTATTATTATTATTATTATTATTATTATTATTATTAT                                                                                                                                                                                                                                                              | ДН≀        |                  | _                                           |                           | н          | •                                       | S              |                 |
| CAGGTGCGTGCGTGCGGGGGGGGGGGGGGGGGGGGGGGG                                                                                                                                                                                                                                                                                                                   | OZI        | CAA              | H                                           | CCI                       | <b>!</b> > | SGA                                     | <b>≯</b> 1     | rat<br>Ata      |
| CAGG<br>GTCC<br>GGAC<br>GGAC<br>CCTG<br>CCTC<br>TAAA                                                                                                                                                                                                                                                                                                      | >          | TG(<br>AC(       | w                                           | AG(                       |            | CT(                                     | ×              | AT              |
|                                                                                                                                                                                                                                                                                                                                                           | $\sim$     | 166<br>200       | H                                           | TG                        | S          | SAG                                     | <u>ں</u>       | rtt<br>\AA      |
|                                                                                                                                                                                                                                                                                                                                                           | U.         | C2<br>G1         |                                             | 55                        | M          | 33                                      | П              | A7<br>T7        |

Figure 5E: V heavy chain 4 (VH4) gene sequence (continued)

|                                 | •                                                           |                  |                                        |                       |                              |
|---------------------------------|-------------------------------------------------------------|------------------|----------------------------------------|-----------------------|------------------------------|
| D T S K N Q F S L K L S<br>Nspv | AAACTGAGCA<br>TTTGACTCGT                                    | r<br>D<br>M      | TTGGGGCGGC<br>AACCCCGCCG               | Q G T L V T V<br>Styl | TGGTGACGGT                   |
| ı                               |                                                             | R H              |                                        | T<br>T                |                              |
| လ                               | AGC                                                         | C A<br>BSSHII    | 000                                    | ტ                     | ~<br>GCA<br>CGI              |
| Ľι                              | ATACTT CGAAAACCA GTTTAGCCTG<br>TATGAA GCTTTTTGGT CAAATCGGAC |                  | ATTGCGCGCG                             | Q<br>StyI             | C CAAGGCACCC<br>G GTTCCGTGGG |
| Ol .                            | CA                                                          | <b>.</b>         | TT                                     | ဗ                     | ပ်ပွဲပ                       |
| Z                               | AAC                                                         | , P4             | GTA                                    | Μ                     | GGG<br>GGG                   |
| × ×                             | GTTGATACTT CGAAAAACCA<br>CAACTATGAA GCTTTTTGGT              | D T A V Y Y Eagi | GATACG GCCGTGTATT<br>CTATGC CGGCACATAA | M D X W G             | TTATTGGGGC<br>AATAACCCCG     |
| S<br>NspV                       | ~ E-€                                                       | គ<br>ខ្ល<br>2    | ២០                                     |                       |                              |
| E                               | ACT<br>rGA                                                  | ⊟∵ì              | GATACG<br>CTATGC                       |                       | GATGGA                       |
| Д                               | AT                                                          | Ω                | GA                                     |                       | GAJ                          |
| >                               | GTTG                                                        | Ø                | ၁၅၁၁<br>၅၁၅၅                           | <b>A</b>              | ATGC                         |
| တ                               |                                                             | A                |                                        | ≯                     |                              |
|                                 | rta(<br>Aat(                                                | E                | ACG(<br>FGC(                           | দি                    | SAAZ                         |
| T I<br>BstEII                   | ~~~~<br>GACCATTAGC<br>CTGGTAATCG                            | >                | GCGTGACGGC<br>CGCACTGCCG               | O<br>O                | GATGGCTTTT<br>CTACCGAAAA     |
| Bst                             | GACC                                                        | ഗ                | )<br>)<br>)<br>)                       | Ω                     | GAT                          |
|                                 |                                                             |                  |                                        |                       |                              |

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S S BlpI TAGCTCAG ATCGAGTC

| တ                                                               | CGGGCGAAAG<br>GCCCGCTTTC                       | M            | AGCTATTGGA<br>TCGATAACCT                      | Н            | GATGGGCATT<br>CTACCCGTAA         |
|-----------------------------------------------------------------|------------------------------------------------|--------------|-----------------------------------------------|--------------|----------------------------------|
| 丘                                                               | GA<br>CT                                       |              | TT.<br>AA                                     | U            | 999                              |
| ម                                                               | )<br>()<br>()                                  | X M          | TA<br>SAT                                     | υ<br>Σ       | 160<br>ACC                       |
|                                                                 | 9<br>0<br>0<br>0                               | S            | \GC                                           | 2,           | 3A1<br>CT <i>I</i>               |
| V K K P                                                         |                                                |              |                                               | M            |                                  |
| ×.                                                              | GTGAAAAAAC<br>CACTTTTTTG                       | H            | TTCCTTTACG<br>AAGGAAATGC                      |              | GTCTCGAGTG<br>CAGAGCTCAC         |
| щ                                                               | AAZ<br>TT                                      | S<br>Н       | TT.<br>AA'                                    | L E<br>XhoI  | ~~~~~~<br>CTCGAG<br>GAGCTC       |
| ×                                                               | AA                                             |              | CT                                            | L<br>Xh      | TC<br>ABG                        |
| >                                                               | TG                                             | Ŋ            | 'TC<br>AAG                                    |              | °TC                              |
|                                                                 | _                                              | <b>.</b> .   |                                               | Ŋ            |                                  |
| 团                                                               | CGGCGCGGAA<br>GCCGCGCCTT                       | У            | GTTCCGGATA<br>CAAGGCCTAT                      | ₹<br>D       | AGC<br>TCC                       |
| G A E                                                           | 300                                            | S G<br>BspEI | ~~~~~<br>TCCGGA<br>AGGCCT                     |              | GA.                              |
| ~                                                               | 0<br>0<br>0<br>0                               | S G<br>BspE. |                                               | ъ<br>С       | ~ ~<br>0<br>0<br>0<br>0<br>0     |
| Ŋ                                                               | Ğ<br>C<br>C                                    |              | TT:                                           | Δ.           | <br>CT<br>GA                     |
| 0.4                                                             |                                                | Ŋ            |                                               | X            | \$ (D, C)                        |
| sequence<br>Q S                                                 | SAG                                            | ·<br>•       | AAG<br>PTC                                    | M<br>BstXI   | ~~~<br>ATC<br>FAC                |
| gene sequence                                                   | ~~<br>TGGTTCAGAG<br>ACCAAGTCTC                 | N<br>N       | AGCTGCAAAG<br>TCGACGTTTC                      | Q M<br>BstXI | ccagarg ccrgg<br>ggrcrac ggacc   |
| ) gene<br>V                                                     | rt(<br>AA(                                     | Ö            | TG(<br>AC(                                    |              | ~ CCC                            |
| (VH5)                                                           | ~<br>GG,                                       | ഗ            |                                               | 以            | 999                              |
| iin S<br>L                                                      | CAAT TG<br>GTTA AC                             |              | K L                                           | _            | <u> </u>                         |
| awy chain<br>Q L<br>MfeI                                        | ~~~<br>XAT                                     | Н            | ATT<br>IAZ                                    |              | 3G]                              |
| heav<br>C                                                       | ~^<br>3C <i>P</i><br>3G1                       | ×.           | AAA<br>TTT                                    | 3            | TG(<br>AC(                       |
| 5F: V                                                           | GT(<br>CA(                                     | L<br>K       | GA                                            | Ŋ            | , D                              |
| Figure 5F: V heavy chain 5 (VH5) $\odot$ E V Q L V $\odot$ MfeI | GAAGTGCAAT TGGTTCAGAG<br>CTTCACGTTA ACCAAGTCTC | 니            | CCTGAAATT AGCTGCAAAG<br>GGACTTTTAA TCGACGTTTC |              | TTGGCTGGGT GCGCCAGATG CCTGGGAAGG |
| Œ                                                               | <u>ი</u> ი                                     |              | 0 0                                           | H            | T 4                              |

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I Y P G D S D ATTTATCCGG GCGATAGCGA

|                                    |                         |        | •        |            | ٠.                    |               |             |
|------------------------------------|-------------------------|--------|----------|------------|-----------------------|---------------|-------------|
|                                    | M                       |        | •        | 'GGZ       |                       | Ŋ             |             |
|                                    | Ø                       |        |          | AAT        | TTA                   | R W G         |             |
|                                    | A D K S I S T A Y L Q W |        |          | CTTCAATGGA | GAAGTTACCT            |               | 5           |
|                                    | <b>&gt;</b> +           |        |          | AT         | TA                    | C A<br>BssHII | <pre></pre> |
|                                    | Æ                       |        |          | CGT        | GTGGCGCATA            | C<br>Bs       | <b>?</b>    |
|                                    | H                       |        |          | CCG        | SGC                   | >-            |             |
|                                    | -                       |        |          | CA         | TTTCGTAATC GTGGCGCATA | SDTAMYYCB     |             |
|                                    | ഗ                       |        |          | <b>FAG</b> | <b>ATC</b>            | <b>~</b>      |             |
|                                    | H                       |        |          | ATT        | TAZ                   | <b>4</b>      |             |
| ned)                               | ഗ                       |        |          | AGC        | TCG                   | -             |             |
| contin                             | ×                       |        |          | AA         | LL                    | <b>:</b>      |             |
| ence (                             | Ω                       |        |          | ATA        | <b>IAT</b>            | Ω             |             |
| e segu                             | ~                       |        | ٠        | CGG        | 3CC                   | ຜ໌            |             |
| 15) gen                            | 7                       |        |          | AGCGCGGATA | TCGCGCCTAT            | A             |             |
| n 5 (∨                             | 0,                      |        |          |            |                       | ~             |             |
| y chai                             | Н                       |        | ,        | ATT        | TA                    | 7             |             |
| V heav                             | H                       | HH     | <b>?</b> | ACC        | TGG                   | <b>H</b>      |             |
| Figure 5F: V heavy chain 5 (VH5) o | >                       | BStEII | ~~~~~    | GGTGACCATT | CCACTGGTAA            | S S L K       |             |
| Figu                               |                         | Д      | ?        | G          | Ö                     | S             |             |

ATTATIGCGC GCGTIGGGGC

ACGGCCATGT

GCAGCCTGAA AGCGAGCGAT CGTCGGACTT TCGCTCGCTA

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CCCTGGTGAC GGGACCACTG CCGGTTCCGT GGCCAAGGCA CCTAATAACC GGATTATTGG AAATACGCTA TTTATGCGAT CCGCTACCGA GGCGATGGCT

V S S BlpI ~~~~~

GGTTAGCTCA G CCAATCGAGT C

Figure 5G: V heavy chain 6 (VH6) gene sequence

H Ø ഗ Д X 口 G Д G ഗ Ø Ø MfeI O

CTGGTGAAAC CGAGCCAAAC GACCACTTIG GCTCGGTTIG GICCACGITA ACGITGICAG ACCAGGCCCG TGGTCCGGGC TGCAACAGTC CAGGTGCAAT

ഗ Z ഗ ഗ ഗ BSPEI ~~~~~ G ഗ Ø ပ ഗ Ы

TAGCGTGAGC AGCAACAGCG TCGTTGTCGC AAAGGCCTCT ATCGCACTCG TTTCCGGAGA ACCTGTGCGA TGGACACGCT CCTGAGCCTG GGACTCGGAC

口 3 团 XhoI ഗ 24 G വ BstXI ഗ Ö . 4 3 Z 3 Ø . K

CAGTCTCCTG GGCGTGGCCT CGAGTGGCTG GTCAGAGGAC CCGCACCGGA GCTCACCGAC GACCTAAGCG CTGGATTCGC CGGCGTGGAA GCCGCACCTT

~~~~~~~~~

AACGATTATG CGGTGAGCGT TTGCTAATAC GCCACTCGCA Ø CAAATGGTAT GTTTACCATA Ζ × TAATAGCATC GGCCGTACCT ATTATCGTAG വ ĸ CCGGCATGGA Е

Q 편 &	CAGTTTAGCC GTCAAATCGG	Y C A BSSHII	TTATTGCGCG	G Q G T Styl	GCCAAGGCAC CGGTTCCGTG	· ·	
S K N SpV.	TTCGAAAAAC AAGCTTTTTG	EagI	CGGCCGTGTA GCCGGCACAT	M Y O	GATTATTGGG CTAATAACCC		
continued)	ACCCGGATAC TGGGCCTATG	P E D	CCGGAAGATA GGCCTTCTAT	Y A M	TTATGCGATG AATACGCTAC	·	
Figure 5G: V heavy chain 6 (VH6) gene sequence (continued) $K  S  R  I  I  N  P$ $BSABI$ $\sim \sim \sim \sim \sim \sim \sim \sim \sim \sim \sim \sim \sim \sim \sim \sim \sim \sim \sim$	GAAAAGCCGG ATTACCATCA CTTTTCGGCC TAATGGTAGT	L Q L N S V T	TGCAACTGAA CAGCGTGACC ACGTTGACT GTCGCACTGG	R W G G D G F BSSHII	CGTTGGGGCG GCGATGGCTT GCAACCCGAA	L V T V S S BlpI	CCTGGTGACG GTTAGCTCAG

- Figure 6: oligonucleotides for gene synthesis
- O1K1 5'- GAATGCATACGCTGATATCCAGATGACCCAGAG-CCCGTCTAGCCTGAGC -3'
  - **O1K2** 5'- CGCTCTGCAGGTAATGGTCACACGATCACCCAC-GCTCGCGCTCAGGCTAGACGGGC -3'
- **O1K3** 5'- GACCATTACCTGCAGAGCGAGCCAGGGCATTAG-CAGCTATCTGGCGTGGTACCAGCAG -3'
- **O1K4** 5'- CTTTGCAAGCTGCTGGCTGCATAAATTAATAGT-TTCGGTGCTTTACCTGGTTTCTGCTGGTACCACGCCAG -3'
- **O1K5** 5'- CAGCCAGCAGCTTGCAAAGCGGGGTCCCGTCCC-GTTTTAGCGGCTCTGGATCCGGCACTGATTTTAC -3'
- **O1K6** 5'- GATAATAGGTCGCAAAGTCTTCAGGTTGCAGGC-TGCTAATGGTCAGGGTAAAATCAGTGCCGGATCC -3'
- **O2K1** 5'- CGATATCGTGATGACCCAGAGCCCACTGAGCCT-GCCAGTGACTCCGGGCGAGCC -3'
- **O2K2** 5'- GCCGTTGCTATGCAGCAGGCTTTGGCTGCTTCT-GCAGCTAATGCTCGCAGGCTCGCCCGGAGTCAC -3'
- **O2K3** 5'- CTGCTGCATAGCAACGGCTATAACTATCTGGAT-TGGTACCTTCAAAAACCAGGTCAAAGCCC -3'
- **O2K4** 5'- CGATCCGGGACCCCACTGGCACGGTTGCTGCCC-AGATAAATTAATAGCTGCGGGCTTTGACCTGGTTTTTG -3'
- **O2K5** 5'- AGTGGGGTCCCGGATCGTTTTAGCGGCTCTGGA-TCCGGCACCGATTTTACCCTGAAAATTAGCCGTGTG -3'
- **O2K6** 5'- CCATGCAATAATACACGCCCACGTCTTCAGCTT-CCACACGCCTAATTTTCAGGG -3'
- O3K1 5'- GAATGCATACGCTGATATCGTGCTGACCCAGAG-CCCGG -3'
- O3K2 5'- CGCTCTGCAGCTCAGGGTCGCACGTTCGCCCGG-AGACAGGCTCAGGGTCGCCGGGCTCTGGGTCAGC -3'
- O3K3 5'- CCCTGAGCTGCAGAGCGAGCCAGAGCGTGAGCA-GCAGCTATCTGGCGTGGTACCAG -3'

- **O3K4** 5'- GCACGGCTGCTCGCGCCATAAATTAATAGACGC-GGTGCTTGACCTGGTTCTGCTGGTACCACGCCAGATAG -3'
- **O3K5** 5'- GCGCGAGCAGCCGTGCAACTGGGGTCCCGGCGC-GTTTTAGCGGCTCTGGATCCGGCACGGATTTTAC -3'
- **O3K6** 5'- GATAATACACCGCAAAGTCTTCAGGTTCCAGGC-TGCTAATGGTCAGGGTAAAATCCGTGCCGGATC -3'
- **O4K1** 5'- GAATGCATACGCTGATATCGTGATGACCCAGAG-CCCGGATAGCCTGGCG -3'
- **O4K2** 5'- GCTTCTGCAGTTAATGGTCGCACGTTCGCCCAG-GCTCACCGCCAGGCTATCCGGGC -3'
- **O4K3** 5'- CGACCATTAACTGCAGAAGCAGCCAGAGCGTGC-TGTATAGCAGCAACAACAAAAACTATCTGGCGTGGTACCAG 3'
- **O4K4** 5'- GATGCCCAATAAATTAATAGTTTCGGCGGCTGA-CCTGGTTCTGCTGGTACCACGCCAGATAG -3'
- **O4K5** 5'- AAACTATTAATTTATTGGGCATCCACCCGTGAA-AGCGGGGTCCCGGATCGTTTTAGCGGCTCTGGATCCGGCAC-3'
- **O4K6** 5'- GATAATACACCGCCACGTCTTCAGCTTGCAGGG-ACGAAATGGTCAGGGTAAAATCAGTGCCGGATCCAGAGCC -3'
- **O1L1** 5'- GAATGCATACGCTCAGAGCGTGCTGACCCAGCC-GCCTTCAGTGAGTGG -3'
- **O1L2** 5'- CAATGTTGCTGCTGCTGCCGCTACACGAGATGG-TCACACGCTGACCTGGTGCGCCACTCACTGAAGGCGGC -3'
- **O1L3** 5'- GGCAGCAGCAGCAACATTGGCAGCAACTATGTG-AGCTGGTACCAGCAGTTGCCCGGGAC -3'
- **O1L4** 5'- CCGGCACGCCTGAGGGACGCTGGTTGTTATCAT-AAATCAGCAGTTTCGGCGCCCGTCCCGGGCAACTGC -3'
- **O1L5** 5'- CCCTCAGGCGTGCCGGATCGTTTTAGCGGATCC-AAAAGCGGCACCAGCGCGAGCCTTGCG -3'

- **O1L6** 5'- CCGCTTCGTCTTCGCTTTGCAGGCCCGTAATCG-CAAGGCTCGCGCTGG -3'
- **O2L1** 5'- GAATGCATACGCTCAGAGCGCACTGACCCAGCC-AGCTTCAGTGAGCGGC -3'
- **O2L2** 5'- CGCTGCTAGTACCCGTACACGAGATGGTAATGC-TCTGACCTGGTGAGCCGCTCACTGAAGCTGG -3'
- **O2L3** 5'- GTACGGGTACTAGCAGCGATGTGGGCGGCTATA-ACTATGTGAGCTGGTACCAGCAGCATCCCGG -3'
- **O2L4** 5'- CGCCTGAGGGACGGTTGCTCACATCATAAATCA-TCAGTTTCGGCGCCCTTCCCGGGATGCTGCTGGTAC -3'
- **O2L5** 5'- CAACCGTCCCTCAGGCGTGAGCAACCGTTTTAG-CGGATCCAAAAGCGGCAACACCGCGAGCC -3'
- **O2L6** 5'- CCGCTTCGTCTTCCGCTTGCAGGCCGCTAATGG-TCAGGCTCGCGGTGTTGCCG -3'
- **O3L1** 5'- GAATGCATACGCTAGCTATGAACTGACCCAGCC-GCCTTCAGTGAGCG -3'
- O3L2 5'- CGCCCAGCGCATCGCCGCTACACGAGATACGCG-CGGTCTGACCTGGTGCAACGCTCACTGAAGGCGGC -3'
- O3L3 5'- GGCGATGCGCTGGGCGATAAATACGCGAGCTGG-TACCAGCAGAAACCCGGGCAGGCGC -3'
- O3L4 5'- GCGTTCCGGGATGCCTGAGGGACGGTCAGAATC-ATCATAAATCACCAGAACTGGCGCCTGCCCGGGTTTC -3'
- O3L5 5'- CAGGCATCCCGGAACGCTTTAGCGGATCCAACA-GCGGCAACACCGCGACCCTGACCATTAGCGG -3'
- O3L6 5'- CCGCTTCGTCTTCCGCCTGAGTGCCGCTAATGG-TCAGGGTC -3'
- O1246H1 5'- GCTCTTCACCCCTGTTACCAAAGCCCAG-GTGCAATTG -3'
- O1AH2 5 ' GGCTTTGCAGCTCACTTTCACGCTGCTGCCCGG-TTTTTTCACTTCCGCGCCAGACTGAACCAATTGCACCTGGGC-TTTG -3 '

- **O1AH3** 5 ' GAAAGTGAGCTGCAAAGCCTCCGGAGGCACTTT-TAGCAGCTATGCGATTAGCTGGGTGCGCCAAGCCCCTGGGCAG GGTC -3 '
- **O1AH4** 5 ' GCCCTGAAACTTCTGCGCGTAGTTCGCCGTGCC-AAAAATCGGAATAATGCCGCCCATCCACTCGAGACCCTGCCC-AGGGGC -3 '
- **O1AH5** 5 ' GCGCAGAAGTTTCAGGGCCGGGTGACCATTACC GCGGATGAAAGCACCAGCACCGCGTATATGGAACTGAGCAGCC TGCG -3 '
- **O1ABH6** 5'- GCGCGCAATAATACACGGCCGTATCTTCGCT-ACGCAGGCTGCTCAGTTCC -3'
- **O1BH2** 5 ' GGCTTTGCAGCTCACTTTCACGCTCGCGCCCGG-TTTTTTCACTTCCGCGCCGCTCTGAACCAATTGCACCTGGGC-TTTG -3'
- **O1BH4** 5 ' GCCCTGAAACTTCTGCGCGTAGTTCGTGCCGCC-GCTATTCGGGTTAATCCAGCCCATCCACTCGAGACCCTGCCCAGGGGC 3 '
- **O1BH5** 5 ' GCGCAGAAGTTTCAGGGCCGGGTGACCATGACC-CGTGATACCAGCATTAGCACCGCGTATATGGAACTGAGCAGCCTGCG -3 '
- **O2H2** 5'- GGTACAGGTCAGGGTCAGGGTTTGGGTCGGTTT-CACCAGGGCCGGCCGCTTTCTTTCAATTGCACCTGGGCTTTG-3'
- **O2H3** 5'- CTGACCCTGACCTGTACCTTTTCCGGATTTAGC-CTGTCCACGTCTGGCGTTGGCGTGGGCTGGATTCGCCAGCCGCCTGGGAAAG -3'
- **O2H4** 5'- GCGTTTTCAGGCTGGTGCTATAATACTTATCAT-CATCCCAATCAATCAGAGCCAGCCACTCGAGGGCTTTCCCAGGCGCTGG -3'

- Figure 6: (continued)
- **O2H5** 5'- GCACCAGCCTGAAAACGCGTCTGACCATTAGCA-AAGATACTTCGAAAAATCAGGTGGTGCTGACTATGACCAACAT GG -3'
- **O2H6** 5'- GCGCGCAATAATAGGTGGCCGTATCCACCGGGT-CCATGTTGGTCATAGTCAGC -3'
- O3H1 5'- CGAAGTGCAATTGGTGGAAAGCGGCGGCCCT-GGTGCAACCGGGCGGCAG -3'
- O3H2 5'- CATAGCTGCTAAAGGTAAATCCGGAGGCCGCCCAGCTCAGACGCAGGCTGCCGCCCGGTTGCAC -3'
- O3H3 5'- GATTTACCTTTAGCAGCTATGCGATGAGCTGGG-TGCGCCAAGCCCCTGGGAAGGGTCTCGAGTGGGTGAG -3'
- O3H4 5'- GGCCTTTCACGCTATCCGCATAATAGGTGCTGC-CGCCGCTACCGCTAATCGCGCTCACCCACTCGAGACCC -3'
- **O3H5** 5'- CGGATAGCGTGAAAGGCCGTTTTACCATTTCAC-GTGATAATTCGAAAAAACACCCTGTATCTGCAAATGAACAG-3'
- **O3H6** 5'- CACGCGCGCAATAATACACGGCCGTATCTTCCG-CACGCAGGCTGTTCATTTGCAGATACAGG -3'
- **04H2** 5'- GGTCAGGCTCAGGGTTTCGCTCGGTTTCACCAG-GCCCGGACCACTTTCTTGCAATTGCACCTGGGCTTTG -3'
- **O4H3** 5'- GAAACCCTGAGCCTGACCTGCACCGTTTCCGGA-GGCAGCATTAGCAGCTATTATTGGAGCTGGATTCGCCAGCCGC-3'
- **O4H4** 5'- GATTATAGTTGGTGCTGCCGCTATAATAAATAT-AGCCAATCCACTCGAGACCCTTCCCAGGCGGCTGGCGAATCCAGG-3'
- **O4H5** 5'- CGGCAGCACCAACTATAATCCGAGCCTGAAAAG-CCGGGTGACCATTAGCGTTGATACTTCGAAAAACCAGTTTAGCCTG -3'
- **O4H6** 5'- GCGCGCAATAATACACGGCCGTATCCGCCGCCG-TCACGCTGCTCAGGTTTCAGGCTAAACTGGTTTTTCG -3'

- **O5H1** 5'- GCTCTTCACCCCTGTTACCAAAGCCGAAGTGCA-ATTG -3'.
- **O5H2** 5'- CCTTTGCAGCTAATTTTCAGGCTTTCGCCCGGT-TTTTTCACTTCCGCGCCGCTCTGAACCAATTGCACTTCGGCTTTGG -3'
- **O5H4** 5'- CGGAGAATAACGGGTATCGCTATCGCCCGGATA-AATAATGCCCATCCACTCGAGACCCTTCCCAGGCATCTGGCGCAC -3'
- **O5H5** 5'- CGATACCCGTTATTCTCCGAGCTTTCAGGGCCA-GGTGACCATTAGCGCGGATAAAAGCATTAGCACCGCGTATCTTC-3'
- **O5H6** 5'- GCGCGCAATAATACATGGCCGTATCGCTCGCTT-TCAGGCTGCTCCATTGAAGATACGCGGTGCTAATG -3'
- **O6H2** 5'- GAAATCGCACAGGTCAGGCTCAGGGTTTGGCTC-GGTTTCACCAGGCCCGGACCAGACTGTTGCAATTGCACCTGG-GCTTTG -3'
- O6H3 5'- GCCTGACCTGTGCGATTTCCGGAGATAGCGTGA-GCAGCAACAGCGCGGCGTGGAACTGGATTCGCCAGTCTCCTGGGCG-3'
- **O6H4** 5'- CACCGCATAATCGTTATACCATTTGCTACGATA-ATAGGTACGGCCCAGCCACTCGAGGCCACGCCCAGGAGACTG-GCG-3'
- **O6H5** 5'- GGTATAACGATTATGCGGTGAGCGTGAAAAGCC-GGATTACCATCAACCCGGATACTTCGAAAAACCAGTTTAGCCTGC -3'
- **O6H6** 5'- GCGCGCAATAATACACGGCCGTATCTTCCGGGG-TCACGCTGTTCAGTTGCAGGCTAAACTGGTTTTTC -3'
- OCLK1 5 ' GGCTGAAGACGTGGGCGTGTATTATTGCCAGCA-GCATTATACCACCCCGCCGACCTTTGGCCAGGGTAC 3 '
  SUBSTITUTE SHEET (RULE 26)

- OCLK2 5'- GCGGAAAAATAAACACGCTCGGAGCAGCCACCG-TACGTTTAATTTCAACTTTCGTACCCTGGCCAAAGGTC -3'
- OCLK3 5'- GAGCGTGTTTATTTTTCCGCCGAGCGATGAACA-ACTGAAAAGCGGCACGGCGAGCGTGTGTGCCTGCTG -3'
- OCLK4 5'- CAGCGCGTTGTCTACTTTCCACTGAACTTTCGC-TTCACGCGGATAAAAGTTGTTCAGCAGGCACACCACGC -3'
- OCLK5 5 ' GAAAGTAGACAACGCGCTGCAAAGCGGCAACAG-CCAGGAAAGCGTGACCGAACAGGATAGCAAAGATAG -3 '
- OCLK6 5 ' GTTTTTCATAATCCGCTTTGCTCAGGGTCAGGG-TGCTGCTCAGAGAATAGGTGCTATCTTTGCTATCCTGTTCG -3 '
- OCLK7 5 ' GCAAAGCGGATTATGAAAAACATAAAGTGTATG CGTGCGAAGTGACCCATCAAGGTCTGAGCAGCCCGGTG 3 '
- OCLK8 5 ' GGCATGCTTATCAGGCCTCGCCACGATTAAAAG-ATTTAGTCACCGGGCTGCTCAGAC -3'
- **OCH1** 5'- GGCGTCTAGAGGCCAAGGCACCCTGGTGACGGT-TAGCTCAGCGTCGAC -3'
- OCH2 5'- GTGCTTTTGCTGCTCGGAGCCAGCGGAAACACG-CTTGGACCTTTGGTCGACGCTGAGCTAACC -3'
- OCH3 5'- CTCCGAGCAGCAAAAGCACCAGCGGCGCACGG-CTGCCCTGGCTGCCTGGTTAAAGATTATTTCC -3'
- **OCH4** 5'- CTGGTCAGCGCCCCGCTGTTCCAGCTCACGGTG-ACTGGTTCCGGGAAATAATCTTTAACCAGGCA -3'
- OCH5 5'- AGCGGGGCGCTGACCAGCGGCGTGCATACCTTT-CCGGCGGTGCTGCAAAGCAGCGGCCTG -3'
- **OCH6** 5'- GTGCCTAAGCTGCTCGGCACGGTCACAACG-CTGCTCAGGCTATACAGGCCGCTGCTTTGCAG -3'
- OCH7 5'- GAGCAGCAGCTTAGGCACTCAGACCTATATTTG-CAACGTGAACCATAAACCGAGCAACACC -3'
- OCH8 5'- GCGCGAATTCGCTTTTCGGTTCCACTTTTTTAT-CCACTTTGGTGTTGCTCGGTTTATGG -3'

Figure 7A: sequence of the synthetic Ck gene segment

Q GCGATGAACA CGCTACTIGI . 回 D S AAAGGCGGCT TTTCCGCCGA Д Д ᄺ GACGAGGCTC GCACAAATAA CGTGTTTATT H H > CTGCTCCGAG S Д Ø Ø GCATGCCACC CGTACGGTGG BsiWI 11111

TTGAAAATAG AACTTTTATC GGACGACTTG CCTGCTGAAC CCGTGCCGCT CGCACCACAC GCGTGGTGTG Ö > GGCACGGCGA Н C TGACTTTTCG ACTGAAAAGC ഗ

N A L Q S G ACAACGCGCT GCAAAGCGGC CGTTTCGCCG ACCTTTCATC TGTTGCGCGA Ω TGGAAAGTAG > 3 CGCGTGAAGC GAAAGTTCAG CTTTCAAGTC × GCGCACTTCG 团 Д

GCACCTATTC CGTGGATAAG വ TCGTTTCTAT AGCAAAGATA X D വ CGAACAGGAT GCTTGTCCTA ഠ AACAGCCAGG AAAGCGTGAC TTTCGCACTG > လ 回 TTGTCGGTCC O ഗ

AAACATAAAG TTTGTATTTC GGATTATGAA CCTAATACTT H Ω Ø ACCCTGACCC TGAGCAAAGC TGGGACTGGG ACTCGTTTCG ഗ Н TCTGAGCAGC AGACTCGTCG ഗ

Figure 7A: sequence of the synthetic Ck gene segment (continued)

×	GGTGACTAAA	CCACTGATTT
H	BAC	TG
	GGTC	CCAC
Д	CCC	999
S	CAG	3TC(
ß	TGAGCAGCCC	ACTCGTCGGG
Н		
ტ	CATCAAGGTC	GTAGTTCCAG
Oi	ICA.	AGT
H		
T \	SACC	TCACTGG
>	GTC	CAC
凹	CGAAGTGACC	GCTI
ပ	TG	SAC
Ø	TGTATGCGTG	ACATACGCAC
×	FA1	ATA
>	H	AC

SFNRGEA\* StuI SphI

TGC CTGATAAGCA GTGGCGAGGC CACCGCTCCG TCTTTTAATC AGAAAATTAG

Figure 78: sequence of the synthetic CH1 gene segment

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BlpI SalI

AGGCTCGTCG TICCGCIGGC ICCGAGCAGC GGTTCGCACA AAGGCGACCG CCAAGCGTGT CGAGTCGCAG CTGGTTTCCA GACCAAAGGT GCTCAGCGTC

CCGACGGACC AATTTCTAAT GGCTGCCTGG TTAAAGATTA TITICGIGGI CGCCGCGIG CCGACGGGAC GGCTGCCCTG K Ø AAAAGCACCA GCGGCGGCAC ტ ტ ഗ ഗ

CTGACCAGCG GGTCAGTGGC ACTCGACCTT GTCGCCCCGC GACTGGTCGC CCAGTCACCG TGAGCTGGAA CAGCGGGCG ָט ഗ Ζ ഗ T > TTTCCCGGAA AAAGGGCCTT

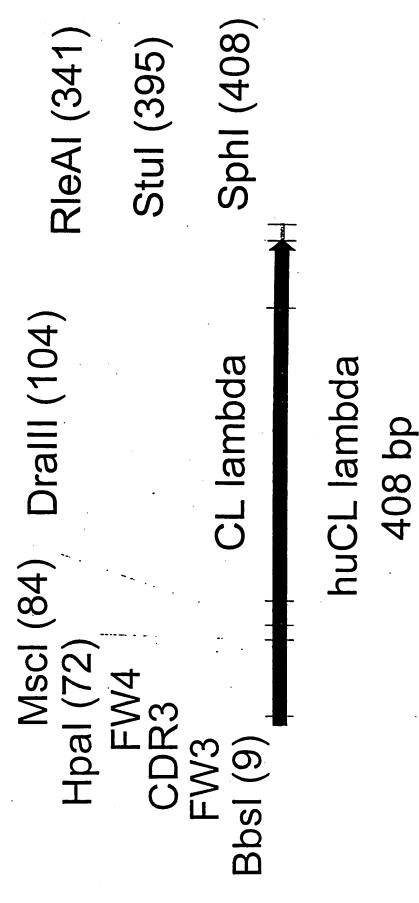
GTATAGCCTG CACGACGTTT CGTCGCCGGA CATATCGGAC ഗ GTGCTGCAAA GCAGCGGCCT ა ტ ഗ U U CGCACGTATG GAAAGGCCGC CTTTCCGGCG Ø Δι GCGTGCATAC

TTAGGCACTC AGACCTATAT AATCCGTGAG TCTGGATATA Ø H ഗ CTCGTCGTCG GAGCAGCAGC ഗ TCGTCGCAAC ACTGGCACGG AGCAGCGTTG TGACCGTGCC ഗ

Figure 7B: sequence of the synthetic CH1 gene segment (continued)

TTTTTTCACC K V D K K V CAAAGTGGAT AAAAAAGTGG GTTTCACCTA N H K P S N T AACCATAAAC CGAGCAACAC TIGGTATITG GCTCGTIGTG TAAGCTT ATTCGAA HindIII · W GCTTAAGACT CGAATTCTGA ECORI ~~~~~ TTGCAACGTG AACGTTGCAC AACCGAAAAG TTGGCTTTTC ഗ × Z Д 回

Figure 7C: functional map and sequence of module 24 comprising the synthetic CA gene segment (huCL lambda)



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Figure 7C: functional map and sequence of module 24 comprising the synthetic CI gene segment (huCL lambda) (continued)

-	BDSL		, C	
⊣	CTTCTGCTTC GCCTAATAAT		CATTATACCA	CCCCGCCTGT
	.i.	HpaI ~~~~~~	MscI	DraIII
51	GTTTGGCGGC GGCACGAAGT TAACCGTTCT CAAACCGCCG CCGTGCTTCA ATTGGCAAGA	GGCACGAAGT TAACCGTTCT CCGTGCTTCA ATTGGCAAGA	TGGCCAGCCG	AAAGCCGCAC TTTCGGCGTG
	DraIII			
101	CGAGTGTGAC GCTGTTTCCG GCTCACACTG CGACAAAGGC	GCTGTTTCCG CCGAGCAGCG CGACAAAGGC GGCTCGTCGC	AAGAATTGCA TTCTTAACGT	GGCGAACAAA CCGCTTGTTT
151	GCGACCCTGG TGTGCCTGAT	TAGCGACTTT	TATCCGGGAG	CCGTGACAGT
	CGCTGGGACC ACACGGACTA	ATCGCTGAAA	ATAGGCCCTC	GGCACTGTCA
201	GGCCTGGAAG GCAGATAGCA	GCCCGTCAA	GGCGGGAGTG	GAGACCACCA
	CCGGACCTIC CGICTATCGI	CGGGGCAGTT	CCGCCCTCAC CTCTGGTGGT	CTCTGGTGGT

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GCCGGTCGTC GATAGACTCG CTATCTGAGC CACCCTCCAA ACAAAGCAAC AACAAGTACG CGGCCAGCAG TTGTTCATGC TGTTTCGTTG GTGGGAGGTT 251

RleAI

**~~~~** 

CTGACGCCTG AGCAGTGGAA GTCCCACAGA AGCTACAGCT GCCAGGTCAC CGGTCCAGTG TCGATGTCGA CAGGGTGTCT TCGTCACCTT GACTGCGGAC

301

StuI

GCATGAGGGG AGCACCGTGG AAAAAACCGT TGCGCCGACT GAGGCCTGAT

CTCCGGACTA TCGTGGCACC TTTTTGGCA ACGCGGCTGA CGTACTCCCC

SphI

?
?
?

401 AAGCATGC TTCGTACG

351

Figure 7D: oligonucleotides used for synthesis of module M24 containing CA gene segment

## M24: assembly PCR

M24-A: GAAGACAAGCGGATTATTATTGCCAGCAGCATTATACCACCCCGCCTGTGTTTGGCGGCG-GCACGAAGTTAACCGTTC

M24-B: CAATTCTTCGCTGCTCGCGGAAACAGCGTCACACTCGGTGCGGCTTTCGGCTGGCCAA-GAACGGTTAACTTCGTGCCGC M24-C: CGCCGAGCAGCGAAGAATTGCAGGCGAACAAAGCGACCCTGGTGTGCCTGATTAGCGACT-TTTATCCGGGAGCCGTGACA

M24-D: TGTTTGGAGGGTGTGGTCTCCACTCCCGCCTTGACGGGGCTGCTATCTGCCTTCCAG-GCCACTGTCACGGCTCCCGG M24-E: CCACACCCTCCAAACAAAGCAACAAGTACGCGGCCAGCAGCTATCTGAGCCTGACGC-CTGAGCAGTGGAAGTCCCACAGAAGCTACAGCTG

M24-F: GCATGCTTATCAGGCCTCAGTCGGCGCAACGGTTTTTTCCACGGTGCTCCCCTCATGCGT-GACCTGGCAGCTGTAGCTTC

CCTAAATGGA AATCGTCGAT ACGCTACTCG ACCCACGCGG TTCGGGGACC

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EH .	ACC IGG	ъ	2 2 2 2 3 2 3	A S Bspei	CCT 3GA	면 건	GCCCCTG
13-VK	T C A	ß	AG	•	900	A P BstXI	i i
agment VHI L SapI	TCTTCACCCC	ഠ	GAAAGCGGCG CTTTCGCCGC	A	CGCGGCCTCC		AAG
ragment V L SapI	}	_		Ö		Q	
thain for L	TTG	>	~ GG1 CC2	ß	GCT	<b>K</b>	CGC
ingle-c	TTACCGTTGC AATGGCAACG	Q MfeI	GCAATTGGTG CGTTAACCAC	. 7	GTCTGAGCTG CAGACTCGAC	V R Q	TGGGTGCGCC AAGCCCCTGG
consensus single	TAC	OE	CAZ	Н	TCT	M	999
conse		>		SLRLSC		·	
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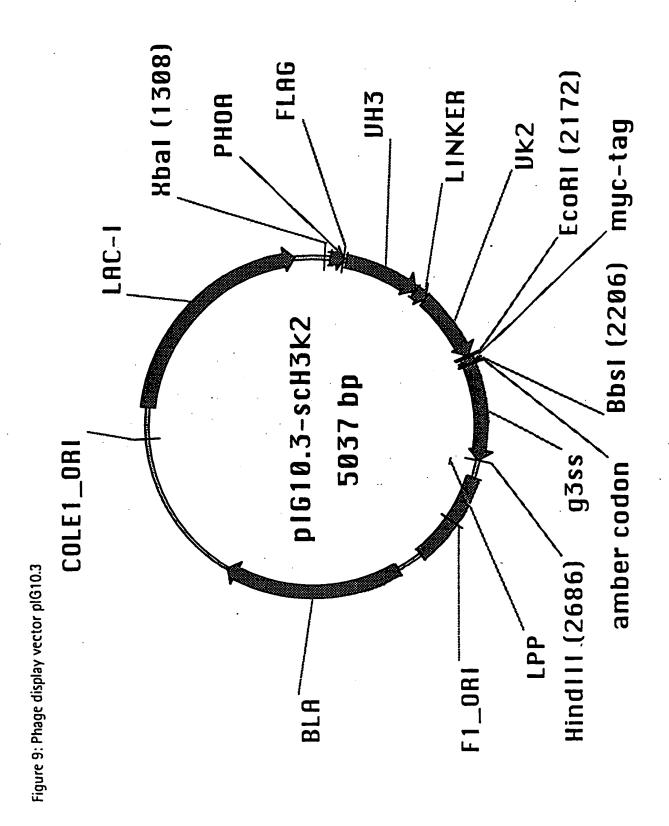
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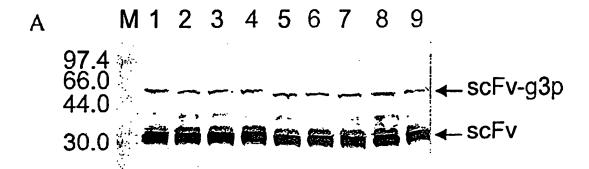
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Figure 11: Expression analysis of initial library



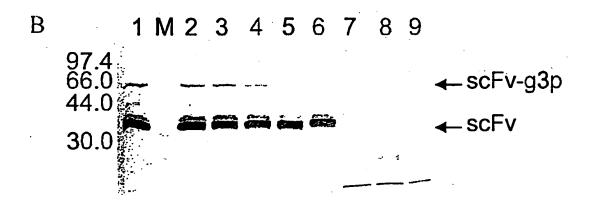


Figure 12: Increase of specificity during the panning rounds

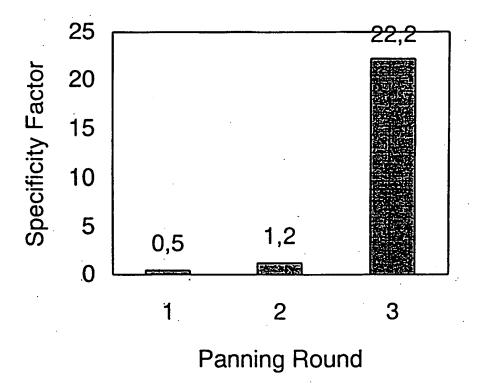


Figure 13: Phage ELISA of clones after the 3rd round of panning

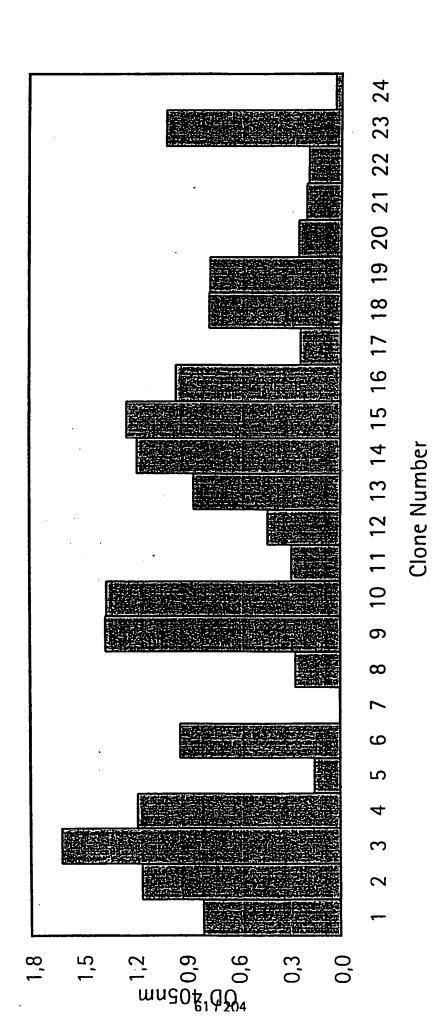
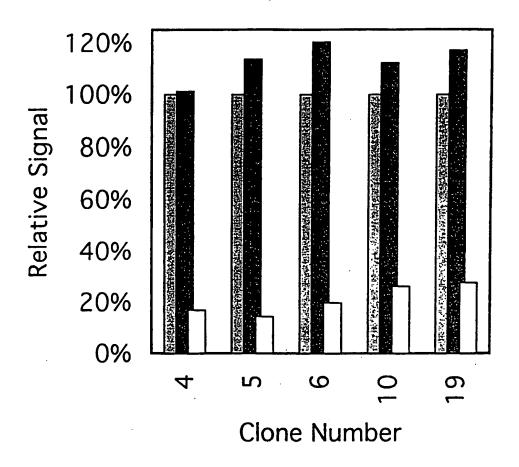


Figure 14: Competition ELISA



- No Inhibition
- Inhibition with BSA
- ☐ Inhibition with Fluorescein

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Figure 16: Purification of fluorescein binding scFv fragments

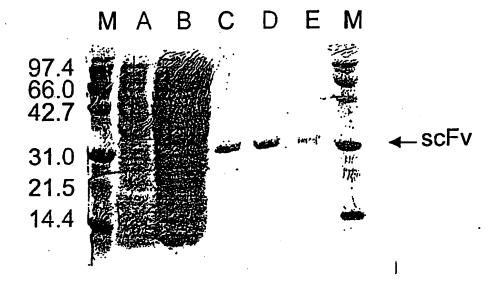
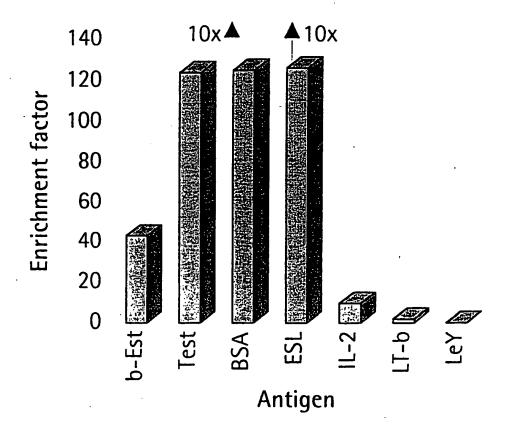


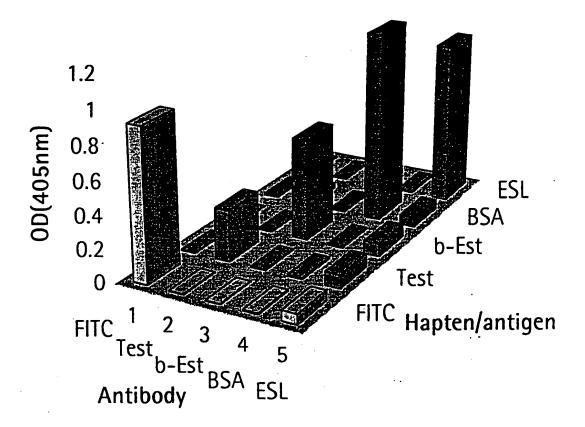
Figure 17: Enrichment factors after three rounds of panning



anti-B-estradiol antibodies *0*9-9 anti-ESL-1 antibodies D9-3 D9-2 PER 1 304 PAR-(mn204)(00 PAR 0

Figure 18: ELISA of anti-ESL-1 and anti- $\beta$ -estradiol antibodies

Figure 19: Selectivity and cross-reactivity of HuCAL antibodies



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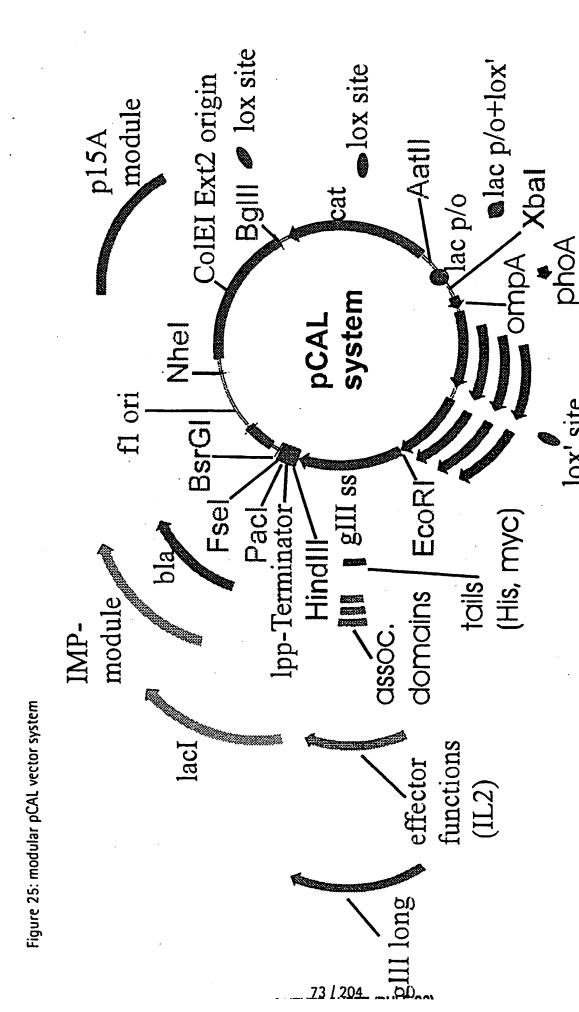


Figure 25a: List of unique restriction sites used in or suitable for HuCAL genes or pCAL vectors

unique restriction site	Isoschizomers
Aatll	
AfIII	Bfrl, BspTl, Bst981
Ascl	
Asel	Vspl, Asnl, PshBl
BamHI	Bstl
Bbel	Ehel, Kasl, Narl
Bbsl	BpuAl, Bpil
BgIII	
Bipi	Bpu1102I,CellI, Blpl
BsaBl	Maml, Bsh1365l, BsrBRl
BsiWl	Pfl23II, SpII, SunI
BspEl	Acclll, BseAl, BsiMl, Kpn2l, Mrol
BsrGl	Bsp1407l, SspBl
BssHII	Paul
BstEII	BstPl, Eco91l, Eco0651
BstXI	
Bsu36l	Aocl, Cvnl, Eco811
Dralll	
DsmAl	
Eagl	BstZl, EclXl, Eco52l, Xmalll
Eco57I	
Eco0109I	Drall
EcoRI	
EcoRV	Eco32l
Fsel	
HindIII	
Hpal	
Kpnl	Acc65l, Asp718l
Mlul	
Mscl	Ball, MluNl

Figure 25a: List of unique restriction sites used in or suitable for HuCAL genes or pCAL vectors

unique restriction site	Isoschizomers
Muni	Mfel
Nhel	1
Nsil	Ppu10l, EcoT22l, Mph1103l
NspV	Bsp119l, BstBl, Csp45l, Lspl, Sful
Pacl	
Pmel	
PmII	BbrPl, Eco72l, PmaCl
Psp5II	PpuMI
Pstl	
Rsrll	(Rsril), Cpol, Cspl
SanDI	
Sapl	
SexAl	
Spel	
Sfil	1
Sphl	Bbul, Pael, Nspl
Stul	Aatl, Eco147l
Styl	Eco130l, EcoT14l
Xbal	BspLU11II
Xhol	PaeR7I
Xmal	Aval, Smal, Cfr9l, PspAl

Figure 26: list of pCAL vector modules

WO 97/08320				PCT/EP96/0364
reference	Skerra et al. (1991) Bio/Technology 9, 273-278	Hoess et al. (1986) Nucleic Acids Res. 2287-2300	see M2	Ge et al., (1994) Expressing antibodies in E. coli. In: Antibody engineering: A practical approach. IRL Press, New York, pp 229-266
template	vector pASK30	(synthetic)	(synthetic)	vector plG10
sites to be inserted	Aatll	lox, BgIII	lox', Sphl	none
sites to be removed	2x Vspl (Asel)	2x Vspl (Asel)	none	Sphl, BamHl
functional element	lac promotor/operator	Cre/lox recombination site	Cre/lox' recombination site	glllp of filamentous phage with N- terminal myctail/amber codon
module/flan- king restriction sites	Aatll-lacp/o- Xbal	BgIII-lox- Aatli	Xbal-lox'- Sphl	EcoRI- gIIIlong- HindIII
ON ON	M L	M2	M3	M7-1

Figure 26: list of pCAL vector modules

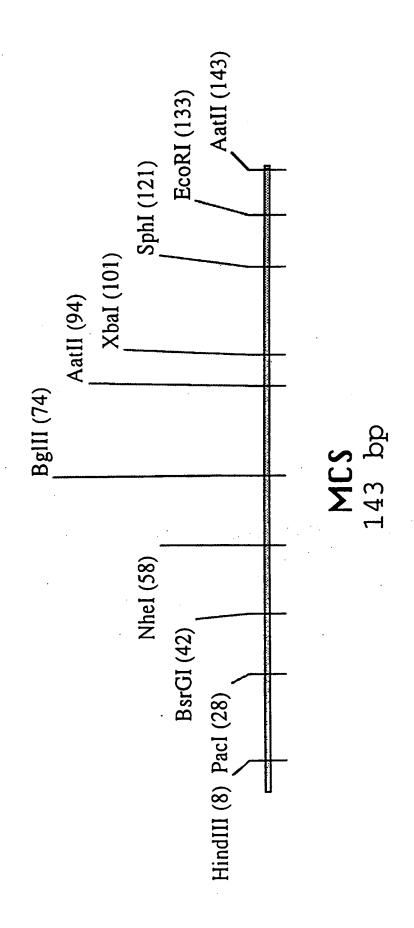
WO 97/08320					PC1/E	P96/0364
see M7-I	see M7-I	see M3	see M1	see M1	see M1	see M1
vector plG10	vector plG10	(synthetic)	(synthetic)	pASK30	pASK30	pASK30
	•	lox	Pacl, Fsel	Pacl, Fsel, BsrGl	BsrGI, Nhel	BsrGI, Nhel
Sphl	Sphl, Bbsl	none	none	Vspl, Eco571, BssSl	Dralll (Banll not removed)	DrallI, BanlI
truncated gillp of filamentous phage with N-terminal Gly- Ser linker	truncated glllp of filamentous phage with N-terminal myctail/amber codon	Cre/lox recombination site	lpp-terminator	beta-lactamase/bla (ampR)	origin of single- stranded replication	origin of single- stranded replication
17-II EcoRI-gillss- HindIII	M7-III EcoRI-gIIIss- HindIII	Sphl-lox- HindIII	HindIII-Ipp- Pacl	Pacl/Fsel-bla- BsrGl	BsrGI-f1 ori- Nhel	BsrGI-f1 ori- Nhel
M7-II	M7-III	M8	M9-11	M10-	M11-	M11-

Figure 26: list of pCAL vector modules

		<del>,</del>			PCT/EP96
Rose, R.E. (1988) Nucleic Acids Res. 16, 355	see M3	Yanisch-Peron, C. (1985) Gene 33,103-119	Cardoso, M. & Schwarz,S. (1992) J. Appl. Bacteriol.72, 289- 293	see M1	Knappik, A & Blückthun, A. (1994) BioTechniques 17, 754-761
pACYC184	(synthetic)	pUC19	pACYC184	(synthetic)	(synthetic)
Nhel, Bglll	BgIII, Iox, Xmnl	BgIII, Nhel			•
BssSI, VspI, NspV	none	Eco571 (BssSl not removed)	BspEl, Mscl, Styl/Ncol	(synthetic)	(synthetic)
origin of double- stranded replication	Cre/lox recombination site	origin of double- stranded replication	chloramphenicol- acetyltransferase/ cat (camR)	signal sequence of phosphatase A	signal sequence of phosphatase A + FLAG detection tag
Nhel-p15A- Bgill	BgIII-lox- BgIII	BgIII-ColEI- Nhel	Aatll-cat- BgIII	Xbal-phoA- EcoRI	Xbal-phoA- FLAG-EcoRI
M12	M13	M14- Ext2	M17	M19	M20

Figure 26: list of pCAL vector modules

WO 97/0832	<u> </u>	
Lee et al. (1983) Infect. Immunol. 264-268	see M1	Lindner et al., (1992) Methods: a companion to methods in enzymology 4, 41- 56
(synthetic)	pASK30	(synthetic)
nal (synthetic)	BstXI, Mlul,BbsI, Banll, BstEII, Hpal, BbeI, VspI	(synthetic)
heat-stable enterotoxin II signal sequence	lac-repressor	poly-histidine tail
Xbal-stll- Sapl	AfIII-laci- Nhel	EcoRI-Histail- HindIII
M21	M41	M42



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Figure 27: functional map and sequence of MCS module (continued)

	HindIII	II:	PacI	BsrGI	
	~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~	<b>?</b>	\ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \	~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~	
Н	ACATGTAAGC	ACATGTAAGC TTCCCCCCCC	CCTTAATTAA GGAATTAATT	CCCCCCCCC TGTACACCCC	
	) 1 1 1 1 1 1				) )
	NheI		Bglii	Aatii Xb	XbaI
	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	<b>?</b>	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	?
51	CCCCCGCTA	CCCCCGCTA GCCCCCCCC	CCAGATCTCC	CCCCCCCGA CGTCCCCCT	,CT
	GGGGGGCGAT	5555555555	CGGGGGGG GGTCTAGAGG	GGGGGGCT GCAGGGGGA	;GA
	XbaI	SphI		EcoRI AatII	
	<b>?</b> ? ? ?	~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~		***************************************	
101	CTAGACCCCC	CCCCCCCATG	ccccccccc	CGAATTCGAC GTC	
	GATCTGGGGG	GGGGCGTAC	9999999999	GCTTAAGCTG CAG	

Figure 28: functional map and sequence of pMCS cloning vector

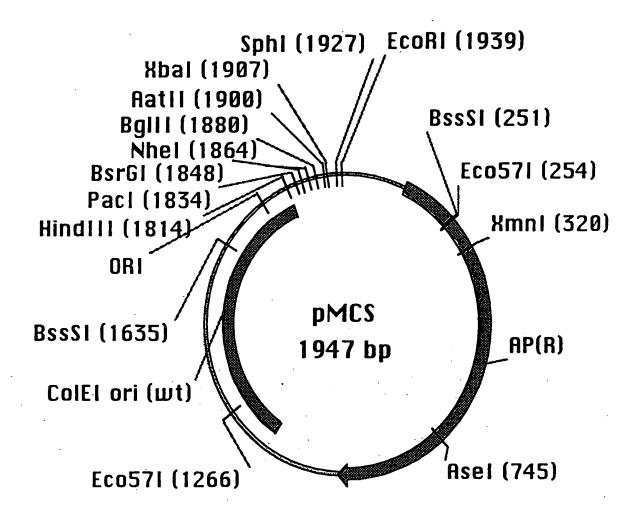


Figure 28: fui	Figure 28: functional map and sequence of pMCS cloning vector (continued)	of pMCS cloning vector (c	continued)		
$\leftarrow$ 1	CAGGTGGCAC GTCCACCGTG	TTTTCGGGGA	AATGTGCGCG TTACACGCGC	GAACCCCTAT CTTGGGGATA	TTGTTTATTT AACAAATAAA
51	TTCTAAATAC AAGATTTATG	ATTCAAATAT TAAGTTTATA	GTATCCGCTC	ATGAGACAAT TACTCTGTTA	AACCCTGATA TTGGGACTAT
101	AATGCTTCAA TTACGAAGTT	TAATATTGAA ATTATAACTT	AAAGGAAGAG TTTCCTTCTC	TATGAGTATT ATACTCATAA	CAACATTTCC GTTGTAAAGG
151	GTGTCGCCCT	TATTCCCTTT ATAAGGGAAA	TTTGCGGCAT	TTTGCCTTCC	TGTTTTTGCT ACAAAAACGA
201	CACCCAGAAA GTGGGTCTTT	CGCTGGTGAA GCGACCACTT	AGTAAAAGAT TCATTTTCTA	Eco57I ~~~~~~ GCTGAAGATC CGACTTCTAG	AGTTGGGTGC TCAACCCACG BSSSI
251	ACGAGTGGGT TGCTCACCCA BssSI	TACATCGAAC ATGTAGCTTG	TGGATCTCAA ACCTAGAGTT	CAGCGGTAAG GTCGCCATTC	ATCCTTGAGA TAGGAACTCT

Figure 28: functional map and sequence of pMCS cloning vector (continued)

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CGTTGCGCAA	ATGGCAACAA	GCCTGTAGCA	ACACCACGAT	GACGAGCGTG	651
CATACCAAAC	TGAATGAAGC	GAACCGGAGC	TGATCGTTGG	TAACTCGCCT	601
GTATGGTTTG	ACTTACTTCG	CTTGGCCTCG	ACTAGCAACC	ATTGAGCGGA	
GGGGATCATG CCCCTAGTAC	GCACAACATG CGTGTTGTAC	CCGCTTTTTT GGCGAAAAAA	AAGGAGCTAA TTCCTCGATT	CGGAGGACCG	551
TGACAACGAT	AACTTACTTC	CACTGCGGCC	TGAGTGATAA	GCCATAACCA	501
ACTGTTGCTA	TTGAATGAAG	GTGACGCCGG	ACTCACTATT	CGGTATTGGT	
ATGCAGTGCT	TAAGAGAATT	GGCATGACAG	TCTTACGGAT	CAGAAAAGCA	451
TACGTCACGA	ATTCTCTTAA	CCGTACTGTC	AGAATGCCTA	GTCTTTTCGT	
TCACCAGTCA AGTGGTCAGT	GGTTGAGTAC CCAACTCATG	AGAATGACTT TCTTACTGAA	CACTATTCTC GTGATAAGAG	TCGCCGCATA	401
AGCAACTCGG TCGTTGAGCC	GCCGGGCAAG CGGCCCGTTC	CCGTATTGAC GGCATAACTG	CGGTATTATC GCCATAATAG	CTATGTGGCG	351
TAAAGTTCTG	TGAGCACTTT	TTTCCAATGA	CGAAGAACGT	GTTTTCGCCC	301
ATTTCAAGAC	ACTCGTGAAA	AAAGGTTACT	GCTTCTTGCA	CAAAAGCGGG	

CATTTTTAAT GTAAAAATTA

TTTAAAACTT AAATTTTGAA

TTTAGATTGA AAATCTAACT

CCAAGTTTAC TCATATATAC GGTTCAAATG AGTATATATG

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Figure 28: functional map and sequence of pMCS cloning vector (continued)

	CTGCTCGCAC	TGTGGTGCTA	CGGACATCGT	TACCGTTGTT	GCAACGCGTT
					AseI
701	ACTATTAACT TGATAATTGA	GGCGAACTAC CCGCTTGATG	TTACTCTAGC AATGAGATCG	TTCCCGGCAA	CAATTAATAG GTTAATTATC
751	ACTGGATGGA TGACCTACCT	GGCGGATAAA CCGCCTATTT	GTTGCAGGAC	CACTTCTGCG GTGAAGACGC	CTCGGCCCTT GAGCCGGGAA
801	CCGGCTGGCT	GGTTTATTGC CCAAATAACG	TGATAAATCT ACTATTTAGA	GGAGCCGGTG CCTCGGCCAC	AGCGTGGGTC TCGCACCCAG
851	TCGCGGTATC AGCGCCATAG	ATTGCAGCAC TAACGTCGTG	TGGGGCCAGA	TGGTAAGCCC ACCATTCGGG	TCCCGTATCG AGGGCATAGC
901	TAGTTATCTA ATCAATAGAT	CACGACGGGG GTGCTGCCCC	AGTCAGGCAA TCAGTCCGTT	CTATGGATGA GATACCTACT	ACGAAATAGA TGCTTTATCT
951	CAGATCGCTG GTCTAGCGAC	AGATAGGTGC TCTATCCACG	CTCACTGATT GAGTGACTAA	AAGCATTGGT TTCGTAACCA	AACTGTCAGA TTGACAGTCT

Figure 28: functional map and sequence of pMCS cloning vector (continued)

1051	1101	1151	1201	1251		1301	1351
H	1	<del></del>	1	1		1	11
TTAAAAGGAT AATTTTCCTA	CCTTAACGTG GGAATTGCAC	CAAAGGATCT GTTTCCTAGA	AAACAAAAAA TTTGTTTTTT	CTACCAACTC GATGGTTGAG	1	AAATACTGTC TTTATGACAG	CTGTAGCACC
CTAGGTGAAG GATCCACTTC	AGTTTTCGTT TCAAAAGCAA	TCTTGAGATC AGAACTCTAG	ACCACCGCTA TGGTGGCGAT	TTTTTCCGAA AAAAAGGCTT		CTTCTAGTGT GAAGATCACA	GCCTACATAC
ATCCTTTTTG	CCACTGAGCG GGTGACTCGC	CTTTTTTCT GAAAAAAAGA	CCAGCGGTGG GGTCGCCACC	GGTAACTGGC CCATTGACCG	{ }	AGCCGTAGTT TCGGCATCAA	CTCGCTCTGC
ATAATCTCAT TATTAGAGTA	TCAGACCCCG	GCGCGTAATC CGCGCATTAG	TTTGTTTGCC AAACAAACGG	C TTCAGCAGAG G AAGTCGTCTC Eco57I	· · · · · · · · · · · · · · · · · · ·	AGGCCACCAC	TAATCCTGTT ATTAGGACAA
GACCAAAATC CTGGTTTTAG	TAGAAAAGAT ATCTTTTCTA	TGCTGCTTGC	GGATCAAGAG CCTAGTTCTC	CGCAGATACC GCGTCTATGG		TTCAAGAACT AAGTTCTTGA	ACCAGTGGCT TGGTCACCGA

	Figure 28: fu	Figure 28: functional map and sequence of pMCS cloning vector (continued)	ce of pMCS cloning vector	(continued)		
	1401	GCTGCCAGTG CGACGGTCAC	GCGATAAGTC CGCTATTCAG	GTGTCTTACC	GGGTTGGACT CCCAACCTGA	CAAGACGATA GTTCTGCTAT
	1451	GTTACCGGAT CAATGGCCTA	AAGGCGCAGC TTCCGCGTCG	GGTCGGGCTG	AACGGGGGGT TTGCCCCCCA	TCGTGCACACACACACACACACACACACACACACGTGTGTGT
	1501	AGCCCAGCTT TCGGGTCGAA	GGAGCGAACG CCTCGCTTGC	ACCTACACCG TGGATGTGGC	AACTGAGATA TTGACTCTAT	CCTACAGCGT GGATGTCGCA
SUBSTITU	1551	GAGCTATGAG	AAAGCGCCAC TTTCGCGGTG	GCTTCCCGAA CGAAGGGCTT	GGGAGAAAGG CCCTCTTTCC	CGGACAGGTA GCCTGTCCAT
ITE SHEET (BUILE 26	1601	TCCGGTAAGC AGGCCATTCG	GGCAGGGTCG	GAACAGGAGA CTTGTCCTCT	GCGCACGAGG CGCGTGCTCC BssSI	GAGCTTCCAG CTCGAAGGTC
3)	1651	GGGGAAACGC CCCCTTTGCG	CTGGTATCTT GACCATAGAA	TATAGTCCTG	TCGGGTTTCG AGCCCAAAGC	CCACCTCTGA
	1701	CTTGAGCGTC GAACTCGCAG	GATTTTTGTG CTAAAAACAC	ATGCTCGTCA	GGGGGGCGGA	GCCTATGGAA CGGATACCTT
	1751	AAACGCCAGC	AACGCGGCCT	TTTTACGGTT	CCTGGCCTTT	TGCTGGCCTT

Figure 28: functional map and sequence of pMCS cloning vector (continued)

ACGACCGGAA	BsrGI ~~~~ CCCCCTGTA GGGGGACAT	AatII ~~~~~ CCCCGACGTC GGGGCTGCAG	EcoRI ~~~~~~~ GAA TTCACGT CTT AAGTGCA
TTGCGCCGGA AAAATGCCAA GGACCGGAAA ACGACCGGAA	PacI ~~~~~~~~ FT AATTAACCCC AA TTAATTGGGG	Bglii ~~~~~~ ccccccag Arcrccccc GGGGGGGTC TAGAGGGGGG	ECORI CCCCCCGAA TTCACGT GGGGGGCTT AAGTGCA
AAAATGCCAA	CCCCCCCTT	Bg CCCCCCCAG GGGGGGGTC	Sphi Accccccc ccareccc Teeeeeee cceraceee
TTGCGCCGGA	HindIII ~~~~~~ GTAAGCTTCC CATTCGAAGG	Nhel ~~~~~~ CCGCTAGCCC GGCGATCGGG	ACCCCCCCC
TTTGCGGTCG	TTGCTCACAT	BsrGI ~~ CACCCCCCC GTGGGGGGG	XbaI ~~~~~~ CCCCCTCTAG ACC GGGGGAGATC TGG
<b>)</b> ,	1801	1851	1901

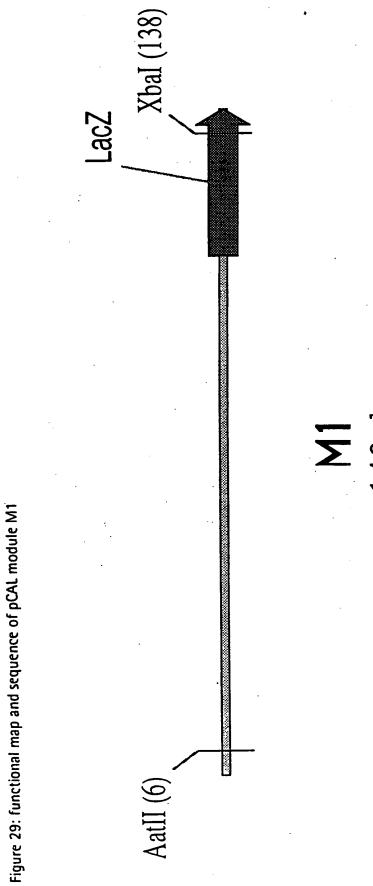


Figure 29: functional map and sequence of pCAL module M1

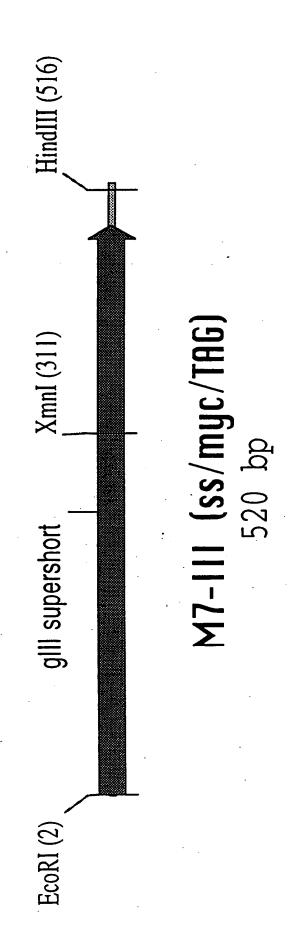
AatII

	\ \ \ \ \				
⊣	GACGTCTTAA	A TGTGAGTTAG	TGTGAGTTAG CTCACTCATT AGGCACCCCA GGCTTTACAC ACACTCAATC GAGTGAGTAA TCCGTGGGGT CCGAAATGTG	AGGCACCCCA TCCGTGGGGT	GGCTTTACAC
51	TTTATGCTTC AAATACGAAG	CGGCTCGTAT GCCGAGCATA	CGGCTCGTAT GTTGTGTGGA ATTGTGAGCG GATAACAATT GCCGAGCATA CAACACCT TAACACTCGC CTATTGTTAA	ATTGTGAGCG TAACACTCGC	GATAACAATT CTATTGTTAA

CGAATTTCTA GCTTAAAGAT AACAGCTATG ACCATGATTA TTGTCGATAC TGGTACTAAT TCACACAGGA 101

XbaI





AATCGGTTGA TTAGCCAACT

TTAATGAATA ATTTCCGTCA ATATTTACCT TCCCTCCTC AATTACTTAT TAAAGGCAGT TATAAATGGA AGGGAGGAG

XmnI

301

Figure 30: functional map and sequence of pCAL module M7-II (continued)

	ECORI				
<del>~</del>	GAATTCGAGC CTTAAGCTCG	AGAAGCTGAT TCTTCGACTA	CTCTGAGGAG GAGACTCCTC	GATCTGTAGG CTAGACATCC	GTGGTGGCTC
51	TGGTTCCGGT ACCAAGGCCA	GATTTTGATT CTAAAACTAA	ATGAAAAGAT TACTTTTCTA	GGCAAACGCT CCGTTTGCGA	AATAAGGGGG TTATTCCCCC
101	CTATGACCGA GATACTGGCT	AAATGCCGAT TTTACGGCTA	GAAAACGCGC	TACAGTCTGA ATGTCAGACT	CGCTAAAGGC GCGATTTCCG
151	AAACTTGATT TTTGAACTAA	CTGTCGCTAC	TGATTACGGT ACTAATGCCA	GCTGCTATCG CGACGATAGC	ATGGTTTCAT
201	TGGTGACGTT ACCACTGCAA	TCCGGCCTTG	CTAATGGTAA GATTACCATT	TGGTGCTACT ACCACGATGA	GGTGATTTTG CCACTAAAAC
251	CTGGCTCTAA GACCGAGATT	TTCCCAAATG AAGGGTTTAC	GCTCAAGTCG CGAGTTCAGC	GTGACGGTGA	TAATTCACCT ATTAAGTGGA

Figure 30: functional map and sequence of pCAL module M7-11 (continued)

TTTTCTATTG AAAAGATAAC	TCTTTTATAT AGAAAATATA	TACTGCGTAA ATGACGCATT		
ACCATATGAA TGGTATACTT	TCTTTGCGTT AGAAACGCAA	TTTGCTAACA TACTGCGTAA AAACGATTGT ATGACGCATT		
GCGCTGGTAA	TTCCGTGGTG	ATTTTCTACG TAAAAGATGC		
TTTGTCTTTG GCGCTGGTAA AAACAGAAAC CGCGACCATT		TTATGTATGT AATACATACA	HindIII	TAAGGAGTCT TGATAAGCTT
ATGTCGCCCT TACAGCGGGA	ATTGTGACAA AATAAACTTA TAACACTGTT TTATTTGAAT	GTTGCCACCT		TAAGGAGTCT
351	401	451		501

Figure 31: functional map and sequence of pCAL module M9-II

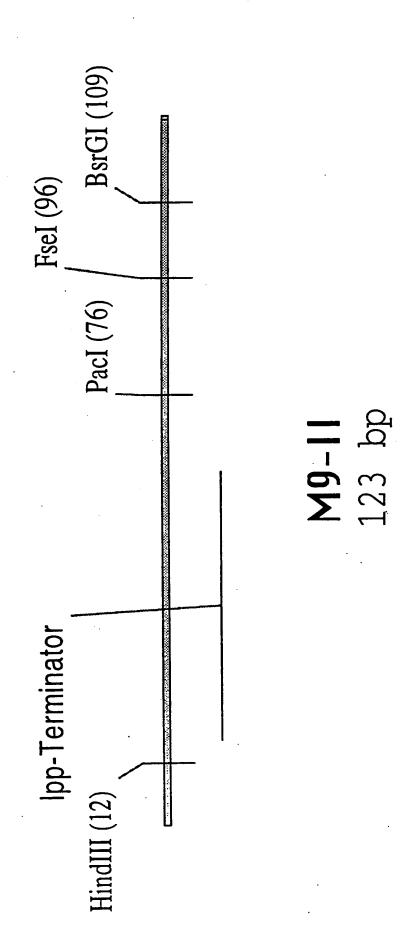


Figure 31: functional map and sequence of pCAL module M9-II (continued)

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	CC TGTGAAGTGA AAAATGGCGC AGATTGTGCG	CGAACTGG ACACTTCACT TTTTACCGCG TCTAACACGC
-	AAAATGGCGC	TTTTACCGCG
-	TGTGAAGTGA	ACACTTCACT
`	AAGCTTGACC	TTCGAACTGG
	9999999999	ವಿರವಿಧಿವಿಧಿವಿಧಿವಿ

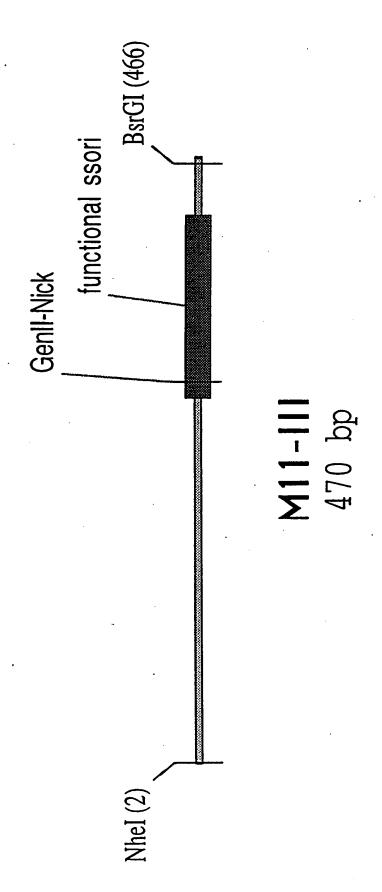
FseI	? ? ? ? ? ? ? ? ? ? ? ? ? ? ? ? ? ? ?	S GGGGGGGGG GCCGGCCTGG	CGGCCGGACC
		5555555555	CCCCCCCC CGGCCGGACC
PacI	~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~	CTGCCGT TTAATTAAAG	GACGCCA AATTAATTTC
		TGTCTGCCGT	ACAGACGGCA
		ACATTTTTT	TGTAAAAAA
		21	

PacI

GGGGGGGTGT ACAGGGGGGG GGG CCCCCCCACA TGTCCCCCCC CCC BsrGI 101

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Figure 32: functional map and sequence of pCAL module M11-III (continued)

		NheI				
	ਜ <b>਼</b>	GCTAGCACGC	GCCCTGTAGC CGGGACATCG	GGCGCATTAA CCGCGTAATT	ວວວອວວອວອວ	TGTGGTGGTT ACACCACCAA
	51	ACGCGCAGCG TGCGCGTCGC	TGACCGCTAC	ACTTGCCAGC TGAACGGTCG	GCCCTAGCGC CGGGATCGCG	CCGCTCCTTT GGCGAGGAAA
•••	101	CGCTTTCTTC	CCTTCCTTTC GGAAGGAAAG	TCGCCACGTT	CGCCGGCTTT GCGGCCGAAA	CCCCGTCAAG GGGGCAGTTC
	151	CTCTAAATCG GAGATTTAGC	GGGCATCCCT CCCGTAGGGA	TTAGGGTTCC AATCCCAAGG	GATTTAGTGC CTAAATCACG	TTTACGGCAC AAATGCCGTG
	201	CTCGACCCCA	AAAAACTTGA TTTTTGAACT	TTAGGGTGAT	GGTTCTCGTA CCAAGAGCAT	GTGGGCCATC
	251	GCCCTGATAG CGGGACTATC	ACGGTTTTTC TGCCAAAAAG	GCCCTTTGAC CGGGAAACTG	GTTGGAGTCC CAACCTCAGG	ACGTTCTTTA TGCAAGAAAT
	301	ATAGTGGACT TATCACCTGA	CTTGTTCCAA GAACAAGGTT	ACTGGAACAA TGACCTTGTT	CACTCAACCC GTGAGTTGGG	TATCTCGGTC ATAGAGCCAG
	351	TATTCTTTTG	ATTTATAAGG	GATTTTGCCG	ATTTCGGCCT	ATTGGTTAAA

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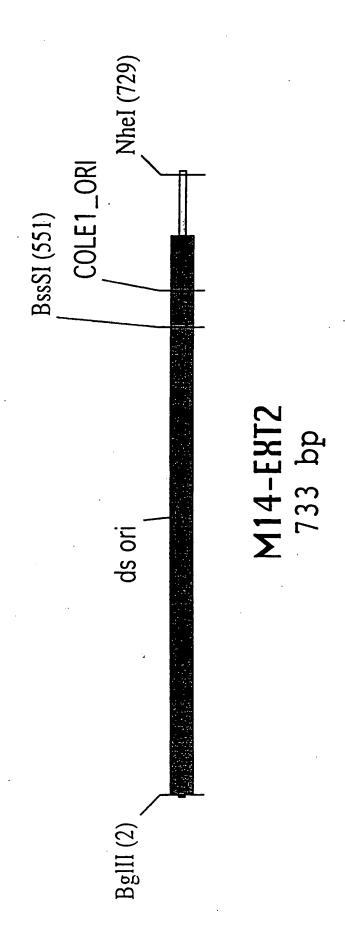
•					
AAAATATTAA TTTTATAATT	ATTTAACAAA AATTTAACGC GAATTTTAAC AAAATATTAA TAAATTGTTT TTAAATTGCG CTTAAAATTG TTTTATAATT	AATTTAACGC TTAAATTGCG	ATTTAACAAA TAAATTGTTT	AAATGAGCTG TTTACTCGAC	401
TAACCAATTT	TAAATATTCC CTAAAACGGC TAAAGCCGGA TAACCAATTT	CTAAAACGGC	TAAATATTCC	ATAAGAAAAC	

BsrGI

CGTTTACAAT TTCATGTACA GCAAATGTTA AAGTACATGT

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Figure 33: functional map and sequence of pCAL module M14-Ext2



GAATGGCCCA

CCAGTGGCGA TAAGTCGTGT CTTACCGGGT GGTCACCGCT ATTCAGCACA GAATGGCCCA

CCTGTTACCA GTGGCTGCTG

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CACCGACGAC

GGACAATGGT

351

ATTCAGCACA

TGGACTCAAG ACGATAGTTA CCGGATAAGG CGCAGCGGTC GGGCTGAACG

GAGACGATTA

TGTATGGAGC

TCGTGGCGGA

TCTTGAGACA

GTGGTGAAGT

Figure 33: functional map and sequence of pCAL module M14-Ext2 (continued)

BglII

	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1				
⊣	AGATCTGACC TCTAGACTGG	AGATCTGACC AAAATCCCTT TCTAGACTGG TTTTAGGGAA	AACGTGAGTT TTGCACTCAA	TTCGTTCCAC	TGAGCGTCAG ACTCGCAGTC
51	ACCCCGTAGA	AAAGATCAAA	GGATCTTCTT	GAGATCCTTT	TTTTCTGCGC
	TGGGGCATCT	TTTCTAGTTT	CCTAGAAGAA	CTCTAGGAAA	AAAAGACGCG
101			AAAAAACCA	SACCHACCAC	ԵԱԱԱԵԵԱԵԵ
H > H	CATTAGACGA	CGAACGTTTG	TTTTTTGGT	GGCGATGGTC	GCCACCAAAC
151	TTTGCCGGAT	CAAGAGCTAC	CAACTCTTTT	TCCGAAGGTA	ACTGGCTACA
	AAACGGCCTA	GTTCTCGATG	GTTGAGAAAA	AGGCTTCCAT	TGACCGATGT
201	GCAGAGCGCA	GATACCAAAT	ACTGTTCTTC	TAGTGTAGCC	GTAGTTAGGC
	CGTCTCGCGT	CTATGGTTTA	TGACAAGAAG	ATCACATCGG	CATCAATCCG
251	CACCACTTCA	CACCACTTCA AGAACTCTGT	AGCACCGCCT	ACATACCTCG CTCTGCTAAT	CTCTGCTAAT

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M14-Ext
module
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map and
are 33: functional
Figure 33:

CCCGACTTGC	ACACCGAACT TGTGGCTTGA	CCCGAAGGGA	AGGAGAGCGC TCCTCGCG BSSSI	GTCCTGTCGG CAGGACAGCC	TCGTCAGGGG	ACGGTTCCTG TGCCAAGGAC
GCGTCGCCAG	CGAACGACCT GCTTGCTGGA	CGCCACGCTT GCGGTGCGAA	GGGTCGGAAC CCCAGCCTTG	TATCTTTATA ATAGAAATAT	TTTGTGATGC	CGGCCTTTTT GCCGGAAAAA
GGCCTATTCC	CAGCTTGGAG GTCGAACCTC	TATGAGAAAG ATACTCTTTC	GTAAGCGGCA CATTCGCCGT	AAACGCCTGG TTTGCGGACC	AGCGTCGATT TCGCAGCTAA	GCCAGCAACG
TGCTATCAAT	GCACACAGCC CGTGTGTCGG	CAGCGTGAGC GTCGCACTCG	CAGGTATCCG GTCCATAGGC	TTCCAGGGGG	CTCTGACTTG	ATGGAAAAAC TACCTTTTTG
Figure 33: Tunctional map and sequence of post models and extensional and sequences of post models and extensional and sequences of post models and extensional and sequences of post models and extensional a	GGGGGTTCGT	GAGATACCTA CTCTATGGAT	GAAAGGCGGA CTTTCCGCCT	ACGAGGGAGC TGCTCCCTCG BssSI	GTTTCGCCAC	GGCGGAGCCT
rigure 33: N	401	451	501	551	601	651

Figure 33: functional map and sequence of pCAL module M14-Ext2 (continued)

NheI

GGCCTTTTGC TCACATGGCT CCGGAAAACG AGTGTACCGA GCCTTTTGCT CGGAAAACGA

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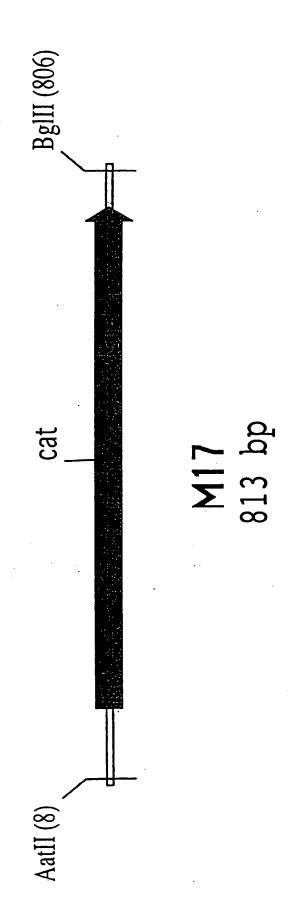


Figure 34: functional map and sequence of pCAL module M17 (continued)

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Н	GGGACGTCGG CCCTGCAGCC	GTGAGGTTCC CACTCCAAGG	AACTTTCACC TTGAAAGTGG	ATAATGAAAT TATTACTTTA	AAGATCACTA TTCTAGTGAT
51	CCGGGCGTAT	TTTTGAGTT AAAAACTCAA	ATCGAGATTT TAGCTCTAAA	TCAGGAGCTA AGTCCTCGAT	AGGAAGCTAA TCCTTCGATT
101	AATGGAGAAA TTACCTCTTT	AAAATCACTG TTTTAGTGAC	GATATACCAC CTATATGGTG	CGTTGATATA GCAACTATAT	TCCCAATGGC AGGGTTACCG
151	ATCGTAAAGA TAGCATTTCT	ACATTTTGAG TGTAAAACTC	GCATTTCAGT	CAGTTGCTCA GTCAACGAGT	ATGTACCTAT TACATGGATA
201	AACCAGACCG TTGGTCTGGC	TTCAGCTGGA	TATTACGGCC ATAATGCCGG	TTTTTAAAGA AAAATTTCT	CCGTAAAGAA GGCATTTCTT
251	AAATAAGCAC TTTATTCGTG	AAGTTTTATC	CGGCCTTTAT	TCACATTCTT AGTGTAAGAA	GCCCGCCTGA CGGGCGGACT
301	TGAATGCTCA ACTTACGAGT	CCCGGAGTTC GGGCCTCAAG	CGTATGGCAA GCATACCGTT	TGAAAGACGG ACTTTCTGCC	TGAGCTGGTG ACTCGACCAC
351	ATATGGGATA	GTGTTCACCC	TTGTTACACC	GTTTTCCATG	AGCAAACTGA

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401	AACGTTTTCA TTGCAAAAGT	TCGCTCTGGA	GTGAATACCA CACTTATGGT	CGACGATTTC GCTGCTAAAG	CGGCAGTTTC GCCGTCAAAG
451	TACACATATA ATGTGTATAT	TTCGCAAGAT AAGCGTTCTA	GTGGCGTGTT CACCGCACAA	ACGGTGAAAA TGCCACTTTT	CCTGGCCTAT GGACCGGATA
501	TTCCCTAAAG AAGGGATTTC	GGTTTATTGA CCAAATAACT	GAATATGTTT CTTATACAAA	TTCGTCTCAG	CCAATCCCTG GGTTAGGGAC
551	GGTGAGTTTC	ACCAGTTTTG TGGTCAAAAC	ATTTAAACGT TAAATTTGCA	AGCCAATATG TCGGTTATAC	GACAACTTCT CTGTTGAAGA
601	TCGCCCCCGT	TTTCACTATG AAAGTGATAC	GGCAAATATT CCGTTTATAA	ATACGCAAGG TATGCGTTCC	CGACAAGGTG GCTGTTCCAC
651	CTGATGCCGC GACTACGGCG	TGGCGATTCA ACCGCTAAGT	GGTTCATCAT CCAAGTAGTA	GCCGTTTGTG CGGCAAACAC	ATGGCTTCCA
701	TGTCGGCAGA	ATGCTTAATG TACGAATTAC	AATTACAACA TTAATGTTGT	GTACTGCGAT CATGACGCTA	GAGTGGCAGG CTCACCGTCC
751	GCGGGGCGTA	ATTTTTTAA	GGCAGTTATT	GGGTGCCCTT	AAACGCCTGG

Figure 34: functional map and sequence of pCAL module M17 (continued)

TAAAAAATT CCGTCAATAA CCCACGGGAA TTTGCGGACC CGCCCCGCAT

BglII

TGCTAGATCT

ACGATCTAGA 801

functional ssori **BsrGI** (612) Hind111 (515) Fse! (599) gill supershort Pac1 (579) GenII-Nick **Kmn1 (310)** Ban II (919) Nhei (1876) replication start ECORI (1) 2755 bp pCAL4 Sph1 (2749) Figure 35: functional map and sequence of modular vector pCAL4 **BssSI (1254)** ColEl Ext2 origin **Kbal** (2739) **Ratil** (2608) lac p/o BgIII (1803) cat

ATCGGTTGAA TAGCCAACTT

TATTTACCTT CCCTCCCTCA

TTTCCGTCAA

AAAGGCAGTT

ATTACTTATT

TAATGAATAA

301

GGGAGGGAGT

ATAAATGGAA

GTGATTTTGC CACTAAAACG

TAATGGTAAT GGTGCTACTG

CCGGCCTTGC

GGTGACGTTT CCACTGCAAA

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GGCCGGAACG

CCACGATGAC

ATTACCATTA

TTAAGTGGAA

ACTGCCACTA

GAGTTCAGCC

AATTCACCTT

CTCAAGTCGG TGACGGTGAT

TCCCAAATGG

TGGCTCTAAT

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ACCGAGATTA

XmnI

Figure 35: functional map and sequence of modular vector pCAL4 (continued)

ECORI

	} } }				
<del></del>	AATTCGAGCA	AATTCGAGCA GAAGCTGATC	TCTGAGGAGG	ATCTGTAGGG	TGGTGGCTCT
: •	TTAAGCTCGT	CTTCGACTAG	AGACTCCTCC	TAGACATCCC	ACCACCGAGA
51		ATTTTGATTA	TGAAAAGATG	GCAAACGCTA	ATAAGGGGGC
	CCAAGGCCAC	ТААААСТААТ	ACTTTTCTAC	CGTTTGCGAT	TATTCCCCCG
101	TATGACCGAA		AAAACGCGCT	ACAGTCTGAC	GCTAAAGGCA
	ATACTGGCTT	TTACGGCTAC	TTTTGCGCGA	TGTCAGACTG	CGATTTCCGT
 151		TGTCGCTACT	GATTACGGTG	CTGCTATCGA	TGGTTTCATT
	TTGAACTAAG	ACAGCGATGA	CTAATGCCAC	GACGATAGCT	ACCAAAGTAA

Figure 35: functional map and sequence of modular vector pCAL4 (continued)

351	TGTCGCCCTT ACAGCGGGAA	TTGTCTTTGG AACAGAAACC	CGCTGGTAAA GCGACCATTT	CCATATGAAT GGTATACTTA	TTTCTATTGA AAAGATAACT
401	TTGTGACAAA AACACTGTTT	ATAAACTTAT TATTTGAATA	TCCGTGGTGT	CTTTGCGTTT GAAACGCAAA	CTTTTATATG GAAAATATAC
451	TTGCCACCTT AACGGTGGAA	TATGTATGTA	TTTTCTACGT	TTGCTAACAT AACGATTGTA	ACTGCGTAAT TGACGCATTA
501	AAGGAGTCTT TTCCTCAGAA	HindIII ~~~~~ GATAAGCTTG CTATTCGAAC	ACCTGTGAAG TGGACACTTC	TGAAAAATGG ACTTTTTACC	CGCAGATTGT
 п п -			Paci		FseI
100	CGCTGTAAAA	AAAACAGACG	GCAAATTAAT	TTCCCCCCC	0000000000
601	TGGGGGGGG	BsrGI ~~~~~~ TGTACATGAA ACATGTACTT	ATTGTAAACG TAACATTTGC	TTAATATTTT AATTATAAAA	GTTAAAATTC CAATTTTAAG

Figure 35: functional map and sequence of modular vector pCAL4 (continued)

AGGCCGAAAT TCCGGCTTTA	GGGTTGAGTG CCCAACTCAC	GGACTCCAAC CCTGAGGTTG	TACGAGAACC ATGCTCTTGG	GCACTAAATC CGTGATTTAG		AAAGCCGGCG TTTCGGCCGC	GCGCTAGGGC CGCGATCCCG
TTTAACCAAT AAATTGGTTA	GACCGAGATA CTGGCTCTAT	TAAAGAACGT ATTTCTTGCA	GATGGCCCAC CTACCGGGTG	GTGCCGTAAA CACGGCATTT		CTTGACGGGG GAACTGCCCC	AAAGGAGCGG TTTCCTCGCC
CAGCTCATTT GTCGAGTAAA	CAAAAGAATA GTTTTCTTAT	AGTCCACTAT TCAGGTGATA	CTATCAGGGC GATAGTCCCG	TGGGGTCGAG		CGATTTAGAG GCTAAATCTC	GAAGAAAGCG CTTCTTTCGC
TTTGTTAAAT AAACAATTTA	CCTTATAAAT GGAATATTTA	TTGGAACAAG	GAAAAACCGT CTTTTTGGCA	TCAAGTTTTT AGTTCAAAAA	BanII		GAAAGGAAGG CTTTCCTTCC
GCGTTAAATT CGCAATTTAA	CGGCAAAATC GCCGTTTTAG	TTGTTCCAGT	GTCAAAGGGC CAGTTTCCCG	ATCACCCTAA TAGTGGGATT		GGAACCCTAA CCTTGGGATT	AACGTGGCGA TTGCACCGCT
651	701	751	801	851		901	951

Figure 35: functional map and sequen 1001 GCTGGCAAGT CGACCGTTCA	1051 TTAATGCGCC AATTACGCGG	1101 AAAAGGCCAG	T 1151 CTCCGCCCCC GAGGGGGGGGGGGGGGGGGGGGGGGGGG	E 1201 GCGAAACCCG	BssSI  1251 CCCTCGTGCG GGGAGCACGC	
Figure 35: functional map and sequence of modular vector pCAL4 (continued) 1001 GCTGGCAAGT GTAGCGGTCA CGCTGC CGTGCCAGT GCGGCAGT GCGACC	GCC GCTACAGGGC CGG CGATGTCCCG	CAG GAACCGTAAA GTC CTTGGCATTT	CCC CTGACGAGCA	CCG ACAGGACTAT	I GCG CTCTCCTGTT ICGC GAGAGGACAA	
.4 (continued) CGCTGCGCGT GCGACGCGCA	NheI ~~~~~~ GCGTGCTAGC CGCACGATCG	AAGGCCGCGT TTCCGGCGCA	TCACAAAAAT AGTGTTTTTA	AAAGATACCA TTTCTATGGT	CCGACCCTGC	
AACCACCACA	CATGTGAGCA GTACACTCGT	TGCTGGCGTT ACGACCGCAA	CGACGCTCAA GCTGCGAGTT	GGCGTTTCCC	CGCTTACCGG	
ອວອວອອວອອອ	AAAGGCCAGC TTTCCGGTCG	TTTCCATAGG	GTCAGAGGTG CAGTCTCCAC	CCTGGAAGCT GGACCTTCGA	ATACCTGTCC TATGGACAGG	C RECECTOR OF A CO

Figure 35: functional map and sequence of modular vector pCAL4 (continued)

	1351	GTATCTCAGT CATAGAGTCA	TCGGTGTAGG	TCGTTCGCTC AGCAAGCGAG	CAAGCTGGGC GTTCGACCCG	TGTGTGCACG ACACACGTGC
	1401	AACCCCCCGT	TCAGCCCGAC AGTCGGGCTG	CGCTGCGCCT GCGACGCGGA	TATCCGGTAA ATAGGCCATT	CTATCGTCTT GATAGCAGAA
9	1451	GAGTCCAACC CTCAGGTTGG	CGGTAAGACA GCCATTCTGT	CGACTTATCG GCTGAATAGC	CCACTGGCAG GGTGACCGTC	CAGCCACTGG
UBSTITUTE	1501	TAACAGGATT ATTGTCCTAA	AGCAGAGCGA TCGTCTCGCT	GGTATGTAGG	CGGTGCTACA GCCACGATGT	GAGTTCTTGA CTCAAGAACT
SHEET (BUI	1551	AGTGGTGGCC TCACCACCGG	TAACTACGGC ATTGATGCCG	TACACTAGAA ATGTGATCTT	GAACAGTATT CTTGTCATAA	TGGTATCTGC ACCATAGACG
E 26)	1601	GCTCTGCTGT	AGCCAGTTAC TCGGTCAATG	CTTCGGAAAA GAAGCCTTTT.	AGAGTTGGTA TCTCAACCAT	GCTCTTGATC
	1651	CGGCAAACAA GCCGTTTGTT	ACCACCGCTG TGGTGGCGAC	GTAGCGGTGG	TTTTTTTGTT AAAAAAACAA	TGCAAGCAGC
	1701	AGATTACGCG TCTAATGCGC	CAGAAAAAAA GTCTTTTTTT	GGATCTCAAG CCTAGAGTTC	AAGATCCTTT TTCTAGGAAA	GATCTTTTCT CTAGAAAAGA

Figure 35: functional map and sequence of modular vector pCAL4 (continued)

1751	3	8	9	9	0	0
5.1	1801	1851	1901	1951	2001	2051
ACGGGGTCTG TGCCCCAGAC	BgllI ~~~~~~ CAGATCTAGC GTCTAGATCG	TACGCCCCGC	TCTGCCGACA	GCGGCATCAG	ACGGGGGCGA	GAAACTCACC CTTTGAGTGG
ACGCTCAGTG TGCGAGTCAC	ACCAGGCGTT TGGTCCGCAA	CCTGCCACTC	TGGAAGCCAT	CACCTTGTCG	AGAAGTTGTC TCTTCAACAG	CAGGGATTGG GTCCCTAACC
GAACGAAAAC CTTGCTTTTG	TAAGGGCACC	ATCGCAGTAC TAGCGTCATG	CACAAACGGC	CCTTGCGTAT	CATATTGGCT GTATAACCGA	CTGAGACGAA GACTCTGCTT
TCACGTTAAG AGTGCAATTC	AATAACTGCC TTATTGACGG	TGTTGTAATT ACAACATTAA	ATGATGAACC TACTACTTGG	AATATTTGCC TTATAAACGG	ACGTTTAAAT TGCAAATTTA	AAACATATTC TTTGTATAAG
GGATTTTGGT CCTAAAACCA	TTAAAAAAT AATTTTTTA	CATTAAGCAT GTAATTCGTA	TGAATCGCCA ACTTAGCGGT	CATAGTGAAA GTATCACTTT	CAAAACTGGT GTTTTGACCA	TCAATAAACC AGTTATTTGG

ATCTTGCGAA TAGAACGCTT	TCCAGAGCGA AGGTCTCGCT	GGGTGAACAC CCCACTTGTG	GAACTCCGGG CTTGAGGCCC	GATAAAACTT CTATTTGAA	TCCAGCTGAA AGGTCGACTT	CTCAAAATGT GAGTTTTACA	CAGTGATTTT GTCACTAAAA
ATCT	TCCA( AGGT(	GGGT	GAAC	GATA CTAT	TCCA AGGT	CTCA	CAGT
AACACGCCAC TTGTGCGGTG	TGGTATTCAC ACCATAAGTG	GGTGTAACAA CCACATTGTT	TTGCCATACG AACGGTATGC	ATAAAGGCCG TATTTCCGGC	GGCCGTAATA	ACTGAAATGC TGACTTTACG	GTGGTATATC CACCATATAG
t (continued) TTTTCACCGT AAAAGTGGCA	GAAATCGTCG CTTTAGCAGC	CATGGAAAAC GTACCTTTTG	CCGTCTTTCA GGCAGAAAGT	AAGAATGTGA TTCTTACACT	TCTTTAAAAA AGAAATTTTT	TGAGCAACTG ACTCGTTGAC	TATATCAACG ATATAGTTGC
e of modular vector pCAL <sup>4</sup> ATAGGCCAGG TATCCGGTCC	GAAACTGCCG CTTTGACGGC	TCAGTTTGCT AGTCAAACGA	CACCAGCTCA	TCAGGCGGGC	TTCTTTACGG AAGAAATGCC	ATAGGTACAT TATCCATGTA	GCCATTGGGA CGGTAACCCT
Figure 35: functional map and sequence of modular vector pCAL4 (continued) 2101 CTTTAGGGAA ATAGGCCAGG TTTTCAGGCAGGTCC AAAAG	TATATGTGTA ATATACACAT	TGAAAACGTT ACTTTTGCAA	TATCCCATAT ATAGGGTATA	TGAGCATTCA	GTGCŤTATTT CACGAATAAA	CGGTCTGGTT	TCTTTACGAT AGAAATGCTA
Figure 35: fi 2 1 0 1	2151	2201	2251	2301	2351	2401	2451
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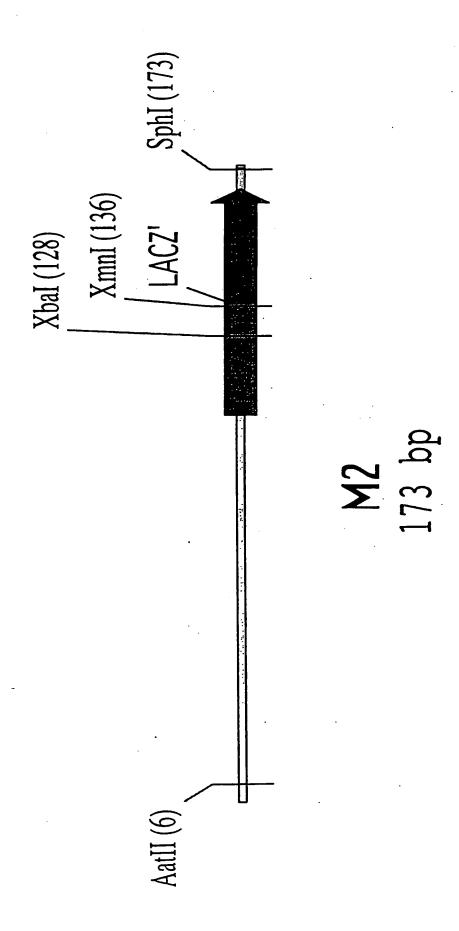


Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

2 Z AatII 111111 GGCTTTACAC CCGAAATGTG AGGCACCCCA TCCGTGGGGT GAGTGAGTAA CTCACTCATT TGTGAGTTAG ACACTCAATC GACGTCTTAA CTGCAGAATT

CTATTGTTAA GATAACAATT ATTGTGAGCG TAACACTCGC GTTGTGTGGA CAACACACCT GCCGAGCATA CGGCTCGTAT AAATACGAAG TTTATGCTTC

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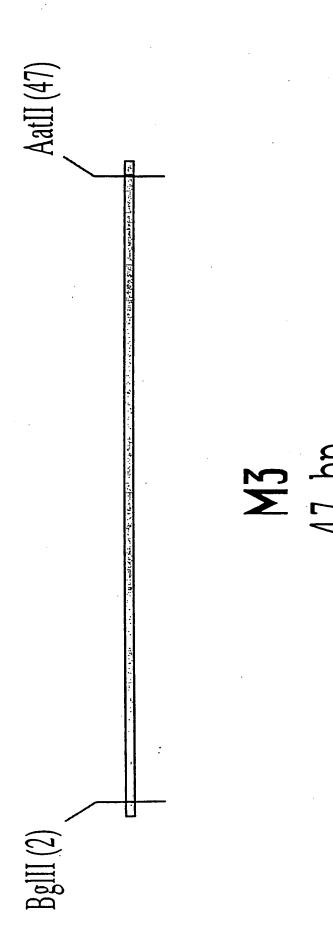
CATATTACAT GTATAATGTA GAATAACTIC CTTATTGAAG ACCATGTCTA TGGTACAGAT AACAGCTATG TTGTCGATAC TCACACAGGA AGTGTGTCCT 101

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 $\mathbf{TGC}$ AGTTATCGCA CGCTATACGA

ACG TCAATAGCGT GCGATATGCT

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AatII

11111

Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

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ACTGCAG TGACGIC ATGCTTCAAT TACGAAGTTA ATGTATGCTA TACATACGAT ACTICGIATA TGAAGCATAT AGATCTCATA TCTAGAGTAT

. Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

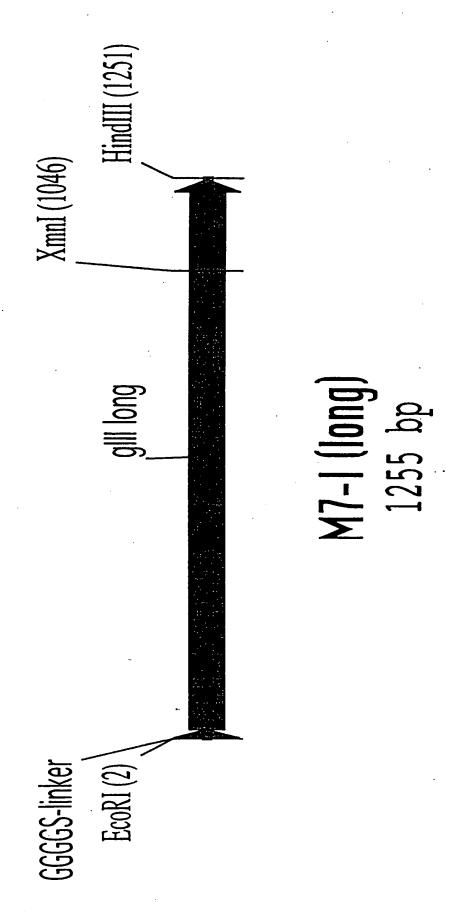


Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

## M 7-I (long):

GAAACGGTTG AAAGTTGTTT CTTTGCCAAC TTTCAACAAA	TAACGTCTGG AAAGACGACA ATTGCAGACC TTTCTGCTGT	GCTGTCTGTG GAATGCTACA CGACAGACAC CTTACGATGT	CAGTGTTACG GTACATGGGT GTCACAATGC CATGTACCCA	GGGTGGTGGC TCTGAGGGTG CCCACCACCG AGACTCCCAC	GCGGTACTAA ACCTCCTGAG CGCCATGATT TGGAGGACTC	TATATCAACC CTCTCGACGG ATATAGTTGG GAGAGCTGCC
GAA	TAA( ATT(	GCT	CAG	300 300 300 300	9090 9090	TAT
TGCGTGCGCT	ATTCATTTAC TAAGTAAATG	AACTATGAGG TTGATACTCC	TGACGAAACT ACTGCTTTGA	CTGAAAATGA GACTTTTACT	TCTGAGGGTG	GGGCTATACT CCCGATATGA
GTGGTGGATC	CATACAGAAA GTATGTCTTT	TCGTTACGCT AGCAATGCGA	TTTGTACTGG AAACATGACC	CTTGCTATCC	GGGTGGCGGT	CACCTATTCC GTGGATAAGG
ECORI ~~~~~~ GAATTCGGTG CTTAAGCCAC	AGCAAAATCC TCGTTTTAGG	AAACTTTAGA TTTGAAATCT	GGCGTTGTAG CCGCAACATC	TCCTATTGGG AGGATAACCC	GCGGTTCTGA CGCCAAGACT	TACGGTGATA
H <sub>.</sub>	51	101	151	201	251	301

Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

351	CACTTATCCG GTGAATAGGC	CCTGGTACTG	AGCAAAACCC TCGTTTTGGG	CGCTAATCCT GCGATTAGGA	AATCCTTCTC TTAGGAAGAG
401	TTGAGGAGTC	TCAGCCTCTT AGTCGGAGAA	AATACTTTCA TTATGAAAGT	TGTTTCAGAA ACAAAGTCTT	TAATAGGTTC ATTATCCAAG
451	CGAAATAGGC GCTTTATCCG	AGGGGGCATT TCCCCCGTAA	AACTGTTTAT TTGACAAATA	ACGGGCACTG TGCCCGTGAC	TTACTCAAGG AATGAGTTCC
501	CACTGACCCC	GTTAAAACTT CAATTTTGAA	ATTACCAGTA TAATGGTCAT	CACTCCTGTA GTGAGGACAT	TCATCAAAAG AGTAGTTTTC
551	CCATGTATGA GGTACATACT	CGCTTACTGG GCGAATGACC	AACGGTAAAT TTGCCATTTA	TCAGAGACTG AGTCTCTGAC	CGCTTTCCAT
601	TCTGGCTTTA	ATGAGGATTT	ATTTGTTTGT TAAACAAACA	GAATATCAAG	GCCAATCGTC CGGTTAGCAG
651	TGACCTGCCT ACTGGACGGA	CAACCTCCTG GTTGGAGGAC	TCAATGCTGG AGTTACGACC	CGGCGGCTCT	GGTGGTGGTT CCACCACCAA
701	CTGGTGGCGG	CTGGTGGCGG CTCTGAGGGT GACCACCGCC GAGACTCCCA	GGTGGCTCTG CCACCGAGAC	AGGGTGGCGG TCCCACCGCC	TTCTGAGGGT AAGACTCCCA

Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

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751	801	851	901	951	1001	1051
GGCGGCTCTG CCGCCGAGAC	TGATTATGAA ACTAATACTT	CCGATGAAAA GGCTACTTTT	GCTACTGATT CGATGACTAA	CCTTGCTAAT GGAACGATTA	AAATGGCTCA TTTACCGAGT	CGTCAATATT GCAGTTATAA
AGGGAGGCGG TCCCTCCGCC	AAGATGGCAA TTCTACCGTT	CGCGCTACAG	ACGGTGCTGC TGCCACGACG	GGTAATGGTG CCATTACCAC	AGTCGGTGAA TCAGCCACTT	TACCTTCCAT
TTCCGGTGGT AAGGCCACCA	ACGCTAATAA TGCGATTATT	TCTGACGCTA AGACTGCGAT	TATCGATGGT ATAGCTACCA	CTACTGGTGA GATGACCACT	GGTGATAATT CCACTATTAA	CCCTCAATCG
GGCTCTGGTT CCGAGACCAA	GGGGCTATG	AAGGCAAACT TTCCGTTTGA	TTCATTGGTG AAGTAACCAC	TTTTGCTGGC AAAACGACCG	CACCTTTAAT GTGGAAATTA	GTTGAATGTC
CCGGTGATTT GGCCACTAAA	ACCGAAAATG TGGCTTTTAC	TGATTCTGTC	ACGTTTCCGG TGCAAAGGCC	TCTAATTCCC AGATTAAGGG	XmnI ~~~~~~~~~ GAATAATTTC CTTATTAAAG	GCCCTTTTGT

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AGTCTTGATA TCAGAACTAT	CGTAATAAGG GCATTATTCC	TAACATACTG	CTACGTTTGC GATGCAAACG	TATGTATTTT	1201
HindIII					
CACCTTTATG GTGGAAATAC	TATATGTTGC ATATACAACG	GGTGTCTTT GCGTTTCTTT CCACAGAAA CGCAAAGAAA	TGGTGTCTTT ACCACAGAAA	ACTTATTCCG TGAATAAGGC	1151
GACAAAATAA CTGTTTTATT	TATTGATTGT GACAAAATAA ATAACTAACA CTGTTTTATT	GGTAAACCCT ATGAATTTTC CCATTTGGGA TACTTAAAAG	GGTAAACCCT CCATTTGGGA	CTTTGGCGCT GAAACCGCGA	1101

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AGCTT TCGAA 1251

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Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

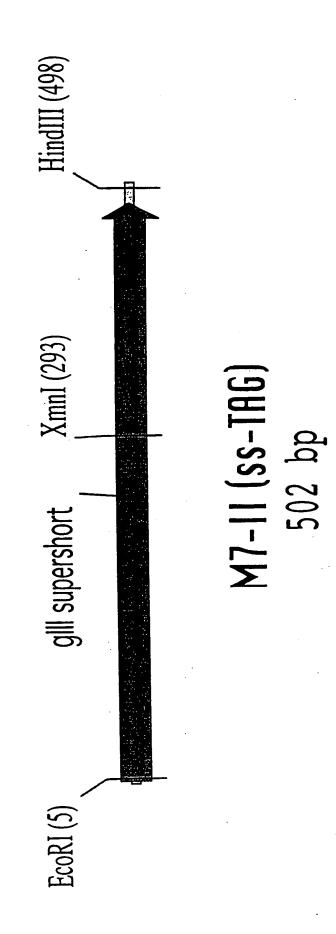


Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

# M 7-II (SS-TAG):

		GTGATTTTGA CACTAAAACT	
		GAGGCGGTTC CGGTGGTGGC TCTGGTTCCG CTCCGCCAAG GCCACCACCG AGACCAAGGC	
		GAGGCGGTTC CGGTGGTGGC TCTGGTTCCG CTCCGCCAAG GCCACCACCG AGACCAAGGC	
ECORI	11111	CGGGAATTCG GCCCTTAAGC	
		Н	

51	TTATGAAAAG AATACTTTTC	ATGGCAAACG CTAATAAGGG TACCGTTTGC GATTATTCCC	CTAATAAGGG GATTATTCCC	ATGGCAAACG CTAATAAGGG GGCTATGACC GAAAATGCCG TACCGTTTGC GATTATTCCC CCGATACTGG CTTTTACGGC	GATATACGGC
101	ATGAAAACGC TACTTTTGCG	GCTACAGTCT	GCTACAGTCT GACGCTAAAG CGATGTCAGA CTGCGATTTC	GCAAACTTGA CGTTTGAACT	TTCTGTCGCT AAGACAGCGA

RECTAT CGATGGTTTC ATTGGTGACG TTTCCGGCCT	IGC AAAGGCCGGA
ATTGGTG1	3 TAACCACTGC
CGATGGTTTC	CGATA GCTACCAAAG TA
GTGCTGCTAT	CACGACGATA
ACTGATTACG	TGACTAATGC
151	

AATTCCCAAA	TTAAGGGTTT
TGCTGGCTCT	ACGACCGAGA
. CTGGTGATTT	GACCACTAAA
AATGGTGCTA	TTACCACGAT
TGCTAATGGT	ACGATTACCA
201	
26)	

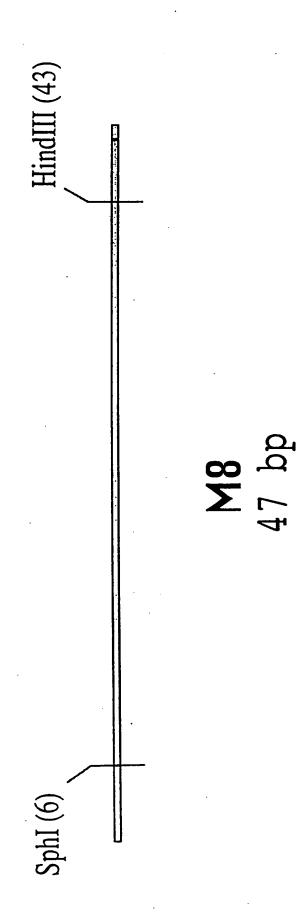
### TAATTTCCGT XmnI CTTTAATGAA GAAATTACTT GATAATTCAC CTATTAAGTG CGGTGACGGT GCCACTGCCA TGGCTCAAGT 251

CTTTTGTCTT GAAAACAGAA	AAAATAAACT TTTTATTTGA	CTTTATGTAT GAAATACATA
GAATGTCGCC CTTACAGCGG	TGATTGTGAC ACTAACACTG	ATGTTGCCAC TACAACGGTG
TCAATCGGTT AGTTAGCCAA	AATTTTCTAT TTAAAAGATA	TGTCTTTGCG TTTCTTTTAT ACAGAAACGC AAAGAAATA
CTTCCCTCCC	AAACCATATG TTTGGTATAC	TGTCTTTGCG
CAATATTTAC GTTATAAATG	TGGCGCTGGT	TATTCCGTGG
301	351	401

CTTGATAAGC GAACTATTCG AATAAGGAGT TTATTCCTCA GTATTTTCTA CGTTTGCTAA CATACTGCGT CATAAAAGAT GCAAACGATT GTATGACGCA 451

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501 TT AA



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Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

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ATGCTTCAAT TACGAAGTTA ATGTACGCTA TACATGCGAT ACTTCGTATA TGAAGCATAT GCATGCCATA CGTACGGTAT

TAAGCTT ATTCGAA

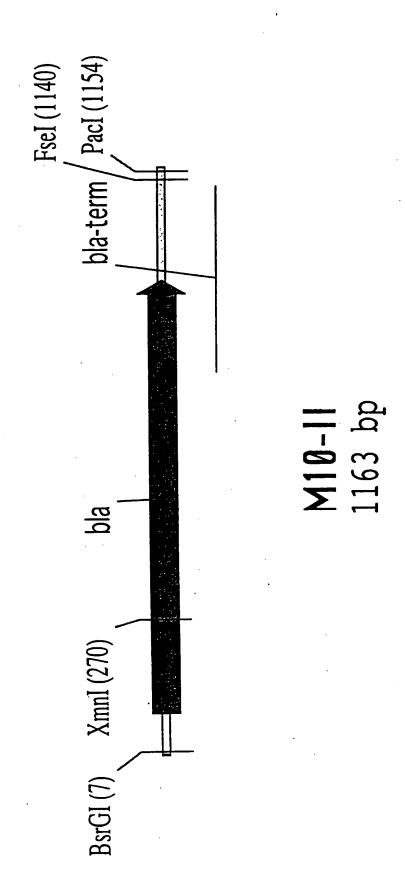


Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

## M 10-II

BsrGI

AACCCTGATA	TTGGGACTAT
ATGAGACAAT	TACTCTGTTA
GTATCCGCTC	CATAGGCGAG
	TAAGTTTATA
GGGGGTGTAC	CCCCCACATG
Н	

CAACATTTCC	GTTGTAAAGG	
TATGAGTATT	ATACTCATAA	
TTGAA AAAGGAAGAG	TTTCCTTCTC	
TAATATTGAA	ATTATAACTT	
AATGCTTCAA	TTACGAAGTT	
21		

TGTTTTGCT	ACAAAAACGA
I TITGCGGCAT TITGCCTTCC	GGGAAA AAACGCCGTA AAACGGAAGG
TTTGCGGCAT	AAACGCCGTA
TATTCCCTTT	SGAA
GTGTCGCCCT	CACAGCGGGA
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AAC TGGATCTCAA CAGCGGTAAG ATCCTTGAGA	3 ACCTAGAGTT GTCGCCATTC TAGGAACTCT
CAGCGGTAAG	GTCGCCATTC
TGGATCTCAA	ACCTAGAGTT
TACATCG	ATGTAGCTTG
GCGAGTGGGT	CGCTCACCCA
201	

AGTTGGGTGC TCAACCCACG

GCTGAGGATC CGACTCCTAG

TCATTTTCTA

CACCCAGAAA CGCTGGTGAA AGTAAAAGAT

GCGACCACTT

GTGGGTCTTT

XmnI

TAAAGTTCTG ATTTCAAGAC TGAGCACTTT ACTCGTGAAA CGAAGAACGT TTTCCAATGA AAAGGTTACT GCTTCTTGCA GTTTTCGCCC CAAAAGCGGG 251

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Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)	COUNTY DATERATION OF A PEASE OF COUNTY BARD
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	401	CAGAAAAGCA GTCTTTTCGT	TCTTACGGAT AGAATGCCTA	GGCATGACAG CCGTACTGTC	TAAGAGAATT ATTCTCTTAA	ATGCAGTGCT
0::5	451	GCCATAACCA CGGTATTGGT	TGAGTGATAA ACTCACTATT	CACTGCGGCC GTGACGCCGG	AACTTACTTC TTGAATGAAG	TGACAACGAT ACTGTTGCTA
	501	CGGAGGACCG GCCTCCTGGC	AAGGAGCTAA TTCCTCGATT	CCGCTTTTTT GGCGAAAAAA	GCACAACATG CGTGTTGTAC	GGGGATCATG CCCCTAGTAC
	551	TAACTCGCCT ATTGAGCGGA	TGATCGTTGG ACTAGCAACC	GAACCGGAGC	TGAATGAAGC ACTTACTTCG	CATACCAAAC GTATGGTTTG
	601	GACGAGCGTG CTGCTCGCAC	ACACCACGAT TGTGGTGCTA	GCCTGTAGCA	ATGGCAACAA TACCGTTGTT	CGTTGCGCAA GCAACGCGTT
	651	ACTATTAACT TGATAATTGA	GGCGAACTAC	TTACTCTAGC AATGAGATCG	TTCCCGGCAA AAGGGCCGTT	CAGTTAATAG GTCAATTATC

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751	CCGGCTGGCT	GGTTTATTGC CCAAATAACG	TGATAAATCT ACTATTTAGA	GGAGCCGGTG	AGCGTGGGTC TCGCACCCAG
801	TCGCGGTATC	ATTGCAGCAC TAACGTCGTG	TGGGGCCAGA	TGGTAAGCCC ACCATTCGGG	TCCCGTATCG AGGGCATAGC
85.1	TAGTTATCTA	CACGACGGGG	AGTCAGGCAA TCAGTCCGTT	CTATGGATGA GATACCTACT	ACGAAATAGA TGCTTTATCT
901	CAGATCGCTG	AGATAGGTGC TCTATCCACG	CTCACTGATT GAGTGACTAA	AAGCATTGGG	TAACTGTCAG ATTGACAGTC
951	ACCAAGTTTA TGGTTCAAAT	CTCATATATA GAGTATATAT	CTTTAGATTG GAAATCTAAC	ATTTAAAACT TAAATTTTGA	TCATTTTTAA AGTAAAAATT
1001	AAATTTTCCT	TCTAGGTGAA	GATCCTTTTT CTAGGAAAAA	GATAATCTCA CTATTAGAGT	TGACCAAAAT ACTGGTTTTA
1051	GGGAATTGCA	GAGTTTTCGT CTCAAAAGCA	TCCACTGAGC AGGTGACTCG	GTCAGACCCC CAGTCTGGGG	GTAGAAAAGA CATCTTTTCT

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Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

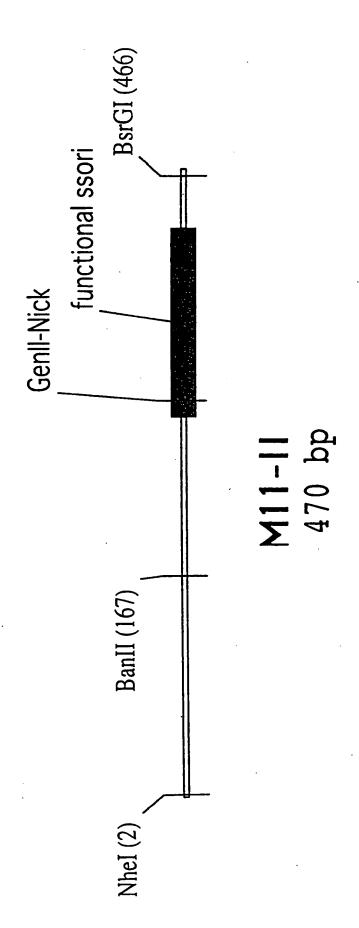


Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

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51	ACGCGCAGCG TGCGCGTCGC	TGACCGCTAC	ACTTGCCAGC TGAACGGTCG	GCCCTAGCGC CGGGATCGCG	CCGCTCCTTT GGCGAGGAAA
101	CGCTTTCTTC GCGAAAGAAG	CCTTCCTTTC GGAAGGAAAG	TCGCCACGTT AGCGGTGCAA	CGCCGGCTTT GCGGCCGAAA	CCCCGTCAAG GGGGCAGTTC
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151	CTCTAAATCG GAGATTTAGC	GGGGCTCCCT CCCCGAGGGA	TTAGGGTTCC AATCCCAAGG	GATTTAGTGC CTAAATCACG	TTTACGGCAC AAATGCCGTG
201	CTCGACCCCA	AAAAACTTGA TTTTTGAACT	TTAGGGTGAT AATCCCACTA	GGTTCTCGTA CCAAGAGCAT	GTGGGCCATC
251	GCCCTGATAG	ACGGTTTTTC TGCCAAAAAG	GCCCTTTGAC	GTTGGAGTCC	ACGTTCTTTA TGCAAGAAAT

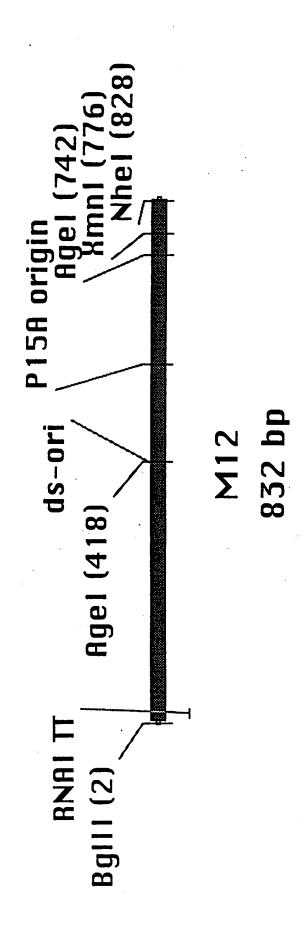
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Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

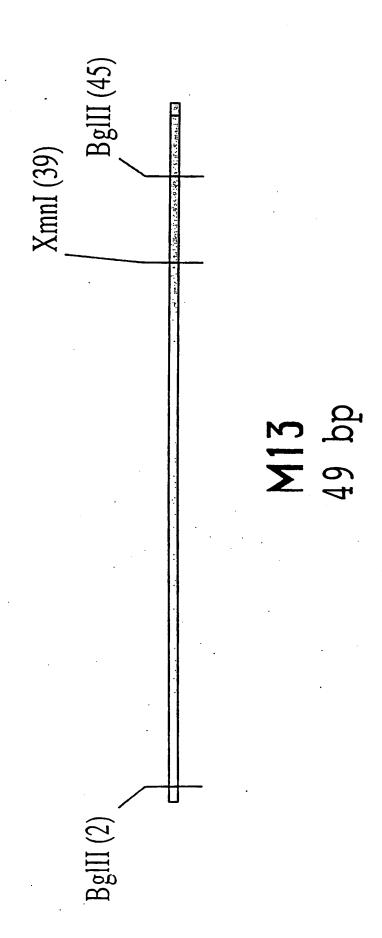


35a: Fur 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued) ${f M-1.2:}$	Bglii ~~~~~~~	1 AGATCTAATA AGATGATCTT CTTGAGATCG TTTTGGTCTG CGCGTAATCT TCTAGATTAT TCTACTAGAA GAACTCTAGC AAAACCAGAC GCGCATTAGA	51 CTTGCTCTGA AAACGAAAAA ACCGCCTTGC AGGGCGGTTT TTCGTAGGTT GAACGAGACT TTTGCTTTTT TGGCGGAACG TCCCGCCAAA AAGCATCCAA	101 CTCTGAGCTA CCAACTCTTT GAACCGAGGT AACTGGCTTG GAGGAGCGCA GAGACTCGAT GGTTGAGAAA CTTGGCTCCA TTGACCGAAC CTCCTCGCGT	151 GTCACTAAAA CTTGTCCTTT CAGTTTAGCC TTAACCGGCG CATGACTTCA CAGTGATTTT GAACAGGAAA GTCAAATCGG AATTGGCCGC GTACTGAAGT	201 AGACTAACTC CTCTAAATCA ATTACCAGTG GCTGCTGCCA GTGGTGCTTT TCTGATTGAG GAGATTTAGT TAATGGTCAC CGACGACGGT CACCACGAAA	251 TGCATGTCTT TCCGGGTTGG ACTCAAGACG ATAGTTACCG GATAAGGCGC ACGTACAGAA AGGCCCAACC TGAGTTCTGC TATCAATGGC CTATTCCGCG	301 AGCGGTCGGA CTGAACGGGG GGTTCGTGCA TACAGTCCAG CTTGGAGCGA TCGCCAGCCT GACTTGCCCC CCAAGCACGT ATGTCAGGTC GAACCTCGCT
(i) <b>2</b> :	igure 3			-	$\vdash$	$\vdash$	7	7	M

Figure 35a 351	Figure 35a: Functional maps and sequences of 351 ACTGCCTACC CGG, TGACGGATGG GCC	. N. E'	additional pCAL vector modules and pCAL vectors (continued) AACTGAG TGTCAGGCGT GGAATGAG, FTGACTC ACAGTCCGCA CCTTACTC	vectors (continued) GGAATGAGAC CCTTACTCTG	AAACGCGGCC TTTGCGCCGG
		AgeI	Į.		
401	ATAACAGCGG TATTGTCGCC	AATGACACCG TTACTGTGGC	GTAAACCGAA CATTTGGCTT	AGGCAGGAAC TCCGTCCTTG	AGGAGAGCGC TCCTCTCGCG
451	AGGAGGGAGC	CGCCAGGGGG	AAACGCCTGG TTTGCGGACC	ТАТСТТТАТА АТАGAAATAT	GTCCTGTCGG
501	GTTTCGCCAC	CACTGATTTG GTGACTAAAC	AGCGTCAGAT TCGCAGTCTA	TTCGTGATGC AAGCACTACG	TTGTCAGGGG
 551	GGCGGAGCCT	ATGGAAAAAC TACCTTTTTG	GGCTTTGCCG CCGAAACGGC	CGGCCCTCTC	ACTTCCCTGT TGAAGGGACA
601	TAAGTATCTT ATTCATAGAA	CCTGGCATCT GGACCGTAGA	TCCAGGAAAT AGGTCCTTTA	CTCCGCCCCG	TTCGTAAGCC
651	ATTTCCGCTC TAAAGGCGAG	GCCGCAGTCG CGGCGTCAGC	AACGACCGAG TTGCTGGCTC	CGTAGCGAGT GCATCGCTCA	CAGTGAGCGA

Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

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CTGCTGACGC	ACTGACACCC	 
ATCACATATT TAGTGTATAA	XmnI ~~~~~~~~~ GAAGCACTTC CTTCGTGAAG	Nhei CAACATAGTA AGCCAGTATA CACTCCGCTA GC GTTGTATCAT TCGGTCATAT GTGAGGCGAT CG
GGAAGCGGAA TATATCCTGT ATCACATATT CCTTCGCCTT ATATAGGACA TAGTGTATAA	CCTTTTTTCT CCTGCCACAT GGAAAAAAGA GGACGGTGTA	Nhe AGCCAGTATA CACTCCGCTA TCGGTCATAT GTGAGGCGAT
GGAAGCGGAA CCTTCGCCTT	CCTTTTTTCT GGAAAAAAGA	CAACATAGTA GTTGTATCAT
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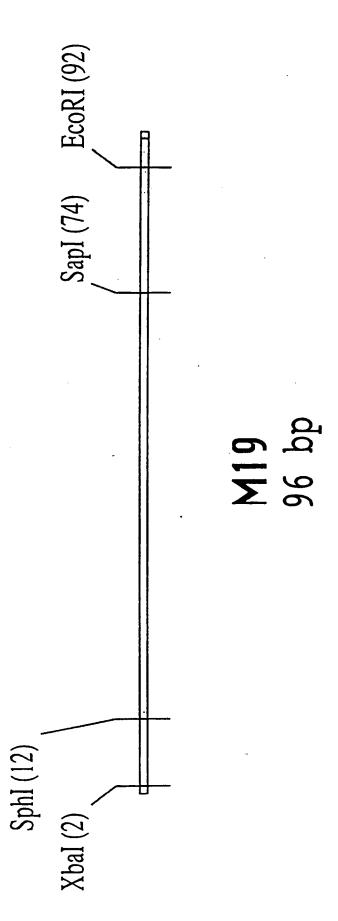
igure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

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AAGTCTAGA TTCAGATCT ATGCTTCAAT TACGAAGTTA ACTICGIATA ATGIATGCIA TGAAGCATAT TACATACGAT AGATCTCATA TCTAGAGTAT





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TTTGTTTCGT

AAACAAAGCA

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ATGGTTTCGG

TACCAAAGCC

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"igure 35a; Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

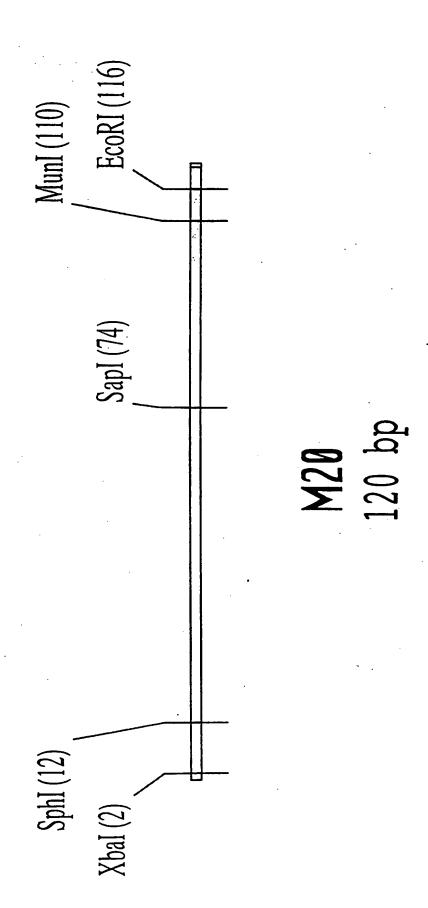
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·	AAATAAAATG TTTATTTTAC	<b>?</b>	TCACCCCTGT AGTGGGGACA	
· · · · · · · · · · · · · · · · · · ·	GCGTAGGAGA CGCATCCTCT	SapI	CCGTTGCTCT GGCAACGAGA	
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Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)



igure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

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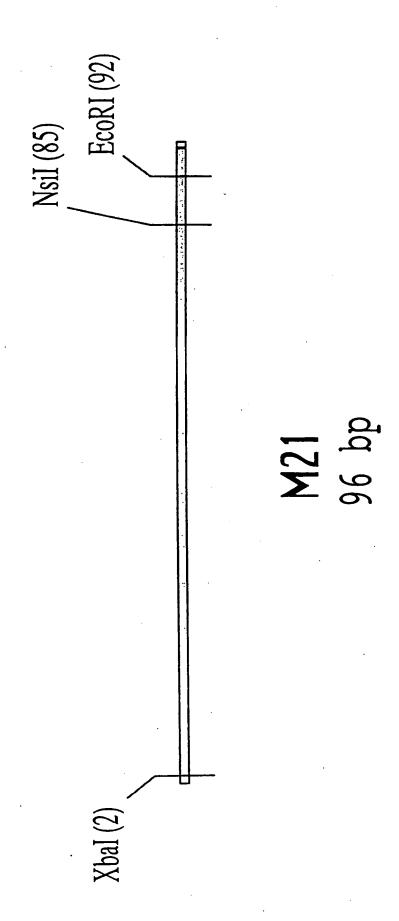
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TATGAAAAAG ATACTTTTTC

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Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

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| 1 1 1 1 | TCTAGAGGTT<br>AGATCTCCAA | • . |
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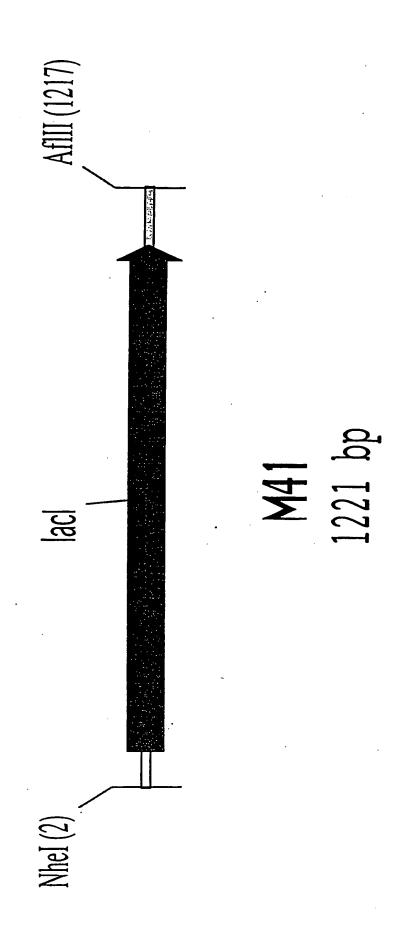


Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

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| Н   | GCTAGCATCG               | AATGGCGCAA<br>TTACCGCGTT | AACCTTTCGC<br>TTGGAAAGCG | GGTATGGCAT<br>CCATACCGTA | GATAGCGCCC<br>CTATCGCGGG |
|-----|--------------------------|--------------------------|--------------------------|--------------------------|--------------------------|
| 51  | GGAAGAGAGT<br>CCTTCTCTCA | CAATTCAGGG               | TGGTGAATGT<br>ACCACTTACA | GAAACCAGTA<br>CTTTGGTCAT | ACGTTATACG<br>TGCAATATGC |
| 101 | ATGTCGCAGA<br>TACAGCGTCT | GTATGCCGGT               | GTCTCTTATC               | AGACCGTTTC<br>TCTGGCAAAG | CCGCGTGGTG<br>GGCGCACCAC |
| 151 | AACCAGGCCA<br>TTGGTCCGGT | GCCACGTTTC<br>CGGTGCAAAG | TGCGAAAACG<br>ACGCTTTTGC | CGGGAAAAAG<br>GCCCTTTTTC | TGGAAGCGGC<br>ACCTTCGCCG |
| 201 | GATGGCGGAG               | CTGAATTACA<br>GACTTAATGT | TTCCTAACCG<br>AAGGATTGGC | CGTGGCACAA<br>GCACCGTGTT | CAACTGGCGG<br>GTTGACCGCC |
| 251 | GCAAACAGTC<br>CGTTTGTCAG | GTTGCTGATT<br>CAACGACTAA | GGCGTTGCCA               | CCTCCAGTCT<br>GGAGGTCAGA | GGCCCTGCAC               |
| 301 | GCGCCGTCGC               | AAATTGTCGC<br>TTTAACAGCG | GGCGATTAAA<br>CCGCTAATTT | TCTCGCGCCG               | ATCAACTGGG<br>TAGTTGACCC |

Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

| GAAGCCTGTA<br>CTTCGGACAT | GCTGATTATT<br>CGACTAATAA                  | CTGCCTGCAC<br>GACGGACGTG | CCCATCAACA<br>GGGTAGTTGT | GGAGCATCTG<br>CCTCGTAGAC | CATTAAGTTC<br>GTAATTCAAG | CTCACTCGCA<br>GAGTGAGCGT | TGCCATGTCC               |
|--------------------------|-------------------------------------------|--------------------------|--------------------------|--------------------------|--------------------------|--------------------------|--------------------------|
| GAAC                     | GCTC                                      | CTG                      | CCC7                     | GGAC                     | CATI                     | CTCA<br>GAG1             | TGCO                     |
| AAGCGGCGTC               | GTGTCAGTGG                                | GCTGTGGAAG               | TGACCAGACA<br>ACTGGTCTGT | GACTGGGCGT<br>CTGACCCGCA | TTAGCTGGCC               | GCATAAATAT<br>CGTATTTATA | GCGACTGGAG               |
| TGGTAGAACG               | CTCGCGCAACGAGGAGGAGGAGGAGGAGGAGGAGGGGGGGG | GGATGCTATT<br>CCTACGATAA | TTGATGTCTC<br>AACTACAGAG | GACGGTACGC               | AATCGCGCTG<br>TTAGCGCGAC | TGGCTGGCTG               | GAACGGGAAG<br>CTTGCCCTTC |
| GTCGTGTCGA               | GCACAATCTT<br>CGTGTTAGAA                  | TGGATGACCA<br>ACCTACTGGT | GCGTTATTTC<br>CGCAATAAAG | CTCCCATGAG<br>GAGGGTACTC | GCCACCAGCA               | CGTCTGCGTC<br>GCAGACGCAG | GCCGATAGCG<br>CGGCTATCGC |
| TGCCAGCGTG<br>ACGGTCGCAC | AAGCGGCGGT<br>TTCGCCGCCA                  | AACTATCCGC<br>TTGATAGGCG | TAATGTTCCG<br>ATTACAAGGC | GTATTATTT<br>CATAATAAAA  | GTCGCATTGG<br>CAGCGTAACC | TGTCTCGGCG               | ATCAAATTCA<br>TAGTTTAAGT |
| 351                      | 401                                       | 451                      | 501                      | 551                      | 601                      | 651                      | 701                      |
|                          |                                           |                          | SUBSTITUT                | E SHEET (R)              | ULE 26)                  |                          |                          |

Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

|             | 751  | GGTTTTCAAC               | AAACCATGCA<br>TTTGGTACGT | AATGCTGAAT<br>TTACGACTTA | GAGGCATCG                | TTCCCACTGC<br>AAGGGTGACG |
|-------------|------|--------------------------|--------------------------|--------------------------|--------------------------|--------------------------|
|             | 801  | GATGCTGGTT<br>CTACGACCAA | GCCAACGATC<br>CGGTTGCTAG | AGATGGCGCT<br>TCTACCGCGA | GGGCGCAATG<br>CCCGCGTTAC | CGTGCCATTA<br>GCACGGTAAT |
|             | 851  | CCGAGTCCGG<br>GGCTCAGGCC | GCTGCGCGTT               | GGTGCGGACA<br>CCACGCCTGT | TCTCGGTAGT               | GGGATACGAC<br>CCCTATGCTG |
| SUBSTITUT   | 901  | GATACCGAGG<br>CTATGGCTCC | ACAGCTCATG<br>TGTCGAGTAC | TTATATCCCG<br>AATATAGGGC | CCGCTGACCA               | CCATCAAACA<br>GGTAGTTTGT |
| E SHEET (RU | 951  | GGATTTTCGC<br>CCTAAAAGCG | CTGCTGGGGC               | AAACCAGCGT<br>TTTGGTCGCA | GGACCGCTTG<br>CCTGGCGAAC | CTGCAACTCT<br>GACGTTGAGA |
| JLE 26)     | 1001 | CTCAGGGCCA               | GGCGGTGAAG<br>CCGCCACTTC | GGCAATCAGC               | TGTTGCCCGT<br>ACAACGGGCA | CTCACTGGTG               |
|             | 1051 | AAAAGAAAAA<br>TTTTCTTTTT | CCACCCTGGC<br>GGTGGGACCG | TCCCAATACG<br>AGGGTTATGC | CAAACCGCCT               | CTCCCGCGC<br>GAGGGGCGCG  |
|             | 1101 | GTTGGCCGAT<br>CAACCGGCTA | TCACTGATGC<br>AGTGACTACG | AGCTGGCACG<br>TCGACCGTGC | ACAGGTTTCC<br>TGTCCAAAGG | CGACTGGAAA<br>GCTGACCTTT |

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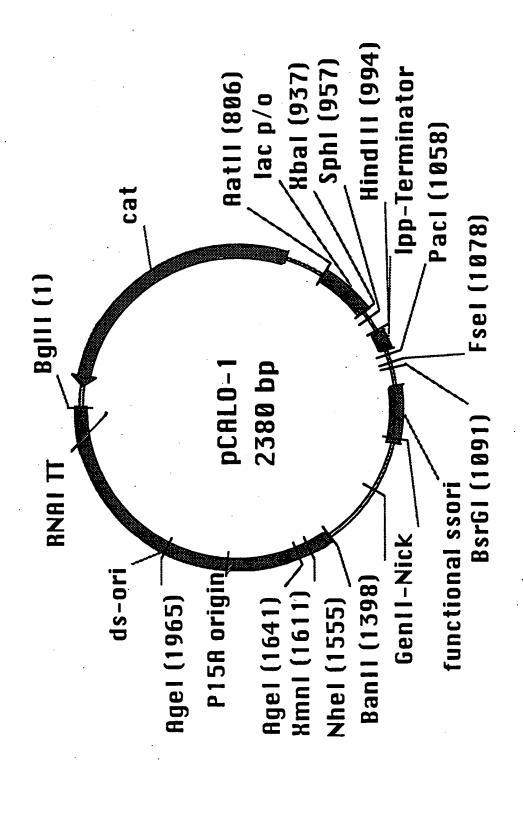
Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

| HATAAAAGCGG CTTCCTGACA GGAGGCCGTT | TATITICGCC GAAGGACTGT CCTCCGGCAA |
|-----------------------------------|----------------------------------|
| ATAAAAG                           | TATTTC                           |
| AGGCTACCCG                        | TCCGATGGGC                       |
| GCGGCCAGTG                        | CGCCCGTCAC                       |
| 1151                              |                                  |

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GCCCACTTAA TTGTTTTGCA AACAAAACGT

Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)



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|-----|------------------------------------------|--------------------------|--------------------------|--------------------------|--------------------------|
| 401 | AAAACGTTTC<br>TTTTGCAAAG                 | AGTTTGCTCA<br>TCAAACGAGT | TGGAAAACGG<br>ACCTTTTGCC | TGTAACAAGG<br>ACATTGTTCC | GTGAACACTA<br>CACTTGTGAT |
| 451 | TCCCATATCA<br>AGGGTATAGT                 | CCAGCTCACC<br>GGTCGAGTGG | GTCTTTCATT               | GCCATACGGA<br>CGGTATGCCT | ACTCCGGGTG<br>TGAGGCCCAC |
| 501 | AGCATTCATC<br>TCGTAAGTAG                 | AGGCGGGCAA               | GAATGTGAAT<br>CTTACACTTA | AAAGGCCGGA<br>TTTCCGGCCT | TAAAACTTGT<br>ATTTTGAACA |
| 551 | GCTTATTTTT<br>CGAATAAAAA                 | CTTTACGGTC<br>GAAATGCCAG | TTTAAAAAGG<br>AAATTTTTCC | CCGTAATATC<br>GGCATTATAG | CAGCTGAACG<br>GTCGACTTGC |
| 601 | GTCTGGTTAT<br>CAGACCAATA                 | AGGTACATTG<br>TCCATGTAAC | AGCAACTGAC<br>TCGTTGACTG | TGAAATGCCT<br>ACTTTACGGA | CAAAATGTTC               |
| 651 | TTTACGATGC                               | CATTGGGATA               | TATCAACGGT               | GGTATATCCA               | GTGATTTTT                |
| 701 |                                          | • •                      | GCTCCTGAAA               | ATCTCGATAA<br>TAGAGCTATT | CTCAAAAAT<br>GAGTTTTTTA  |

| (continued)                                                                                  |
|----------------------------------------------------------------------------------------------|
| L vectors                                                                                    |
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| r modules and                                                                                |
| vecto                                                                                        |
| pCAL                                                                                         |
| Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued) |
| Figure 35a: Fun                                                                              |

| AACCTCACCC<br>TTGGAGTGGG | GCTTTACACT<br>CGAAATGTGA                    | ATAACAATTT<br>TATTGTTAAA |      | ACCCCCCCCC               | HindIII<br>~~~~~~<br>ATAAGCTTGA<br>TATTCGAACT | TTTGTCTGCC               |
|--------------------------|---------------------------------------------|--------------------------|------|--------------------------|-----------------------------------------------|--------------------------|
| TGAAAGTTGG<br>ACTTTCAACC | GGCACCCCAG                                  | TTGTGAGCGG               | Xbal | GAATTTCTAG               | ATACGAAGTT<br>TATGCTTCAA                      | CGACATTTTT<br>GCTGTAAAAA |
| TTCATTATGG<br>AAGTAATACC | TCACTCATTA<br>AGTGAGTAAT                    | TTGTGTGGAA               |      | CCATGATTAC<br>GGTACTAATG | AATGTACGCT<br>TTACATGCGA                      | GCAGATTGTG               |
| GTGATCTTAT<br>CACTAGAATA | GTGAGTTAGC<br>CACTCAATCG                    | GGCTCGTATG               | •    | ACAGCTATGA<br>TGTCGATACT | AACTTCGTAT<br>TTGAAGCATA                      | GAAAAATGGC<br>CTTTTTACCG |
| ACGCCCGGTA               | Aatii<br>~~~~~~<br>GACGTCTAAT<br>CTGCAGATTA | TTATGCTTCC<br>AATACGAAGG |      | CACACAGGAA<br>GTGTGTCCTT | Sphi<br>~~~~~~<br>CGCATGCCAT<br>GCGTACGGTA    | CCTGTGAAGT               |
| 751                      | 801                                         | 851                      |      | 901                      | 951                                           | 1001                     |

Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

| FseI BsrGI | GGGCCGGCCT GGGGGGGGT GTACATGAAA<br>CCCGGCCGGA CCCCCCCA CATGTACTTT | TTAAAATTCG CGTTAAATTT TTGTTAAATC<br>AATTTTAAGC GCAATTTAAA AACAATTTAG | GGCCGAAATC GGCAAAATCC CTTATAAATC<br>CCGGCTTTAG CCGTTTTAGG GAATATTTAG | GGTTGAGTGT TGTTCCAGTT TGGAACAAGA<br>CCAACTCACA ACAAGGTCAA ACCTTGTTCT | GACTCCAACG TCAAAGGGCG AAAAACCGTC<br>CTGAGGTTGC AGTTTCCCGC TTTTTGGCAG | ACGAGAACCA TCACCCTAAT CAAGTTTTTT<br>TGCTCTTGGT AGTGGGATTA GTTCAAAAA | Banli<br>  |
|------------|-------------------------------------------------------------------|----------------------------------------------------------------------|----------------------------------------------------------------------|----------------------------------------------------------------------|----------------------------------------------------------------------|---------------------------------------------------------------------|------------|
|            | AGGGGGGGGG                                                        | TAATATTTTG<br>ATTATAAAAC                                             | TTAACCAATA                                                           | ACCGAGATAG<br>TGGCTCTATC                                             | AAAGAACGTG                                                           | ATGGCCCACT                                                          | ( E        |
| PacI       | 1051 GTTTAATTAA<br>CAAATTAATT                                     | 1101 TTGTAAACGT<br>AACATTTGCA                                        | 1151 AGCTCATTTT<br>TCGAGTAAAA                                        | 1201 AAAAGAATAG<br>TTTTCTTATC                                        | 1251 GTCCACTATT<br>CAGGTGATAA                                        | 1301 TATCAGGGG<br>ATAGTCCCGC                                        | 7547745555 |

Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

| AAAGGAAGGG<br>TTTCCTTCCC | TAGCGGTCAC<br>ATCGCCAGTG | CTACAGGGCG<br>GATGTCCCGC | GATGAGGGTG                                | AgeI    | CCGGTGCGTC   | CACTGACTCG<br>GTGACTGAGC | ACGAACGGGG |
|--------------------------|--------------------------|--------------------------|-------------------------------------------|---------|--------------|--------------------------|------------|
| ACGTGGCGAG A             | CTGGCAAGTG 1             | TAATGCGCCG (ATTACGCGCGC) | TGTTGGCACT (ACAACCGTGA)                   | 7       | AAAGGCTGCA C | CTTCCTCGCT (GAAGGAGCGA   | GAAATGGCTT |
| AAGCCGGCGA<br>TTCGGCCGCT | CGCTAGGGCG               | CCGCCGCGCT               | TGGCTTACTA                                |         | GCAGGAGAAA   | ATATATTCCG<br>TATATAAGGC | GCGGCGAGCG |
| TTGACGGGGA               | AAGGAGCGGG               | ACCACCACAC<br>TGGTGGTGTG | GAGTGTATAC<br>CTCACATATG                  | Н       | GCTTCATGTG   | GTGATACAGG               | TCGTTCGACT |
| GATTTAGAGC<br>CTAAATCTCG | AAGAAAGCGA<br>TTCTTTCGCT | GCTGCGCGTA               | NheI<br>~~~~~<br>CGTGCTAGCG<br>GCACGATCGC | IcmX    | TCAGTGAAGT   | AGCAGAATAT<br>TCGTCTTATA | CTACGCTCGG |
| 1401                     | 1451                     | 1501                     | 1551                                      |         | 1601         | 1651                     | 1701       |
|                          |                          | 5                        | SUBSTITUTE SHEET                          | (RULE 2 | 6)           | •                        |            |

| TGCTTGCCCC                                                                                               | GAAGTGAGAG<br>CTTCACTCTC | GACAAGCATC<br>CTGTTCGTAG | AGGACTATAA<br>TCCTGATATT | CTCCTGTTCC<br>GAGGACAAGG | CGTTTGTCTC                                 | CCAAGCTGGA<br>GGTTCGACCT | TTATCCGGTA<br>AATAGGCCAT |
|----------------------------------------------------------------------------------------------------------|--------------------------|--------------------------|--------------------------|--------------------------|--------------------------------------------|--------------------------|--------------------------|
| TACCGAA                                                                                                  | ACTTAACAGG<br>TGAATTGTCC | CCGCCCCCCT               | GAAACCCGAC<br>CTTTGGGCTG | CTCCTGCGCT               | GTTATGGCCG                                 | GCAGTTCGCT<br>CGTCAAGCGA | CCGCTGCGCC               |
| ules and pCAL vectors (con                                                                               | CCAGGAAGAT<br>GGTCCTTCTA | TCCATAGGCT<br>AGGTATCCGA | CAGTGGTGGC<br>GTCACCACCG | TGGCGGCTCC               | TCATTCCGCT                                 | TTCCGGGTAG               | TTCAGTCCGA               |
| ditional pCAL vector modu<br>AGCAAGCTGA                                                                  | CTGGAAGATG<br>GACCTTCTAC | AAGCCGTTTT<br>TTCGGCAAAA | ACGCTCAAAT<br>TGCGAGTTTA | CGTTTCCCCC               | Agel<br>~~~~~~<br>TTTACCGGTG<br>AAATGGCCAC | TGACACTCAG<br>ACTGTGAGTC | GAACCCCCCG               |
| Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued) | CGGAGATTTC<br>GCCTCTAAAG | GGCCGCGGCA               | ACGAAATCTG<br>TGCTTTAGAC | AGATACCAGG<br>TCTATGGTCC | TGCCTTTCGG                                 | ATTCCACGCC<br>TAAGGTGCGG | CTGTATGCAC<br>GACATACGTG |
| 5a: Functional                                                                                           | 1751                     | 1801                     | 1851                     | 1901                     | 1951                                       | 2001                     | 2051                     |
| Figure 3                                                                                                 |                          |                          | SU                       | 3STITUTE SI:<br>161 / 2  | EET (RULE 26)<br>204                       |                          |                          |

Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

| 2101 ACTATCGTCT TGAGTCCAAC CCG<br>TGATAGCAGA ACTCAGGTTG GGC | 2151 GCAGCCACTG GTAATTGATT TAG<br>CGTCGGTGAC CATTAACTAA ATC | 2201 GTTAAGGCTA AACTGAAAGG<br>CAATTCCGAT TTGACTTTCC | 2251 CAGTTACCTC GGTTCAAAGA GTT<br>GTCAATGGAG CCAAGTTTCT CAA | 2301 GCCCTGCAAG GCGGTTTTTT<br>CGGGACGTTC CGCCAAAAA | E 06) | 2351 CAAAACGATC TCAAGAAGAT CAT |
|-------------------------------------------------------------|-------------------------------------------------------------|-----------------------------------------------------|-------------------------------------------------------------|----------------------------------------------------|-------|--------------------------------|
| CCGGAAAGAC<br>GGCCTTTCTG                                    | _                                                           | ACAAGTTTTA                                          | GTTGGTAGCT CAACCATCGA                                       | CGTTTTCAGA CCAAAAGTCT                              | Bglii | CATCTTATTA                     |
| ATGCAAAAGC<br>TACGTTTTCG                                    | TAGAGGAGTT AGTCTTGAAG<br>ATCTCCTCAA TCAGAACTTC              | GTGACTGCGC                                          | CAGAGAACCT<br>GTCTCTTGGA                                    | GCAAGAGATT<br>CGTTCTCTAA                           |       |                                |
| ACCACTGGCA<br>TGGTGACCGT                                    | TCATGCGCCG                                                  | TCCTCCAAGC                                          | ACGAAAAACC<br>TGCTTTTTGG                                    | ACGCGCAGAC<br>TGCGCGTCTG                           |       |                                |

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Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

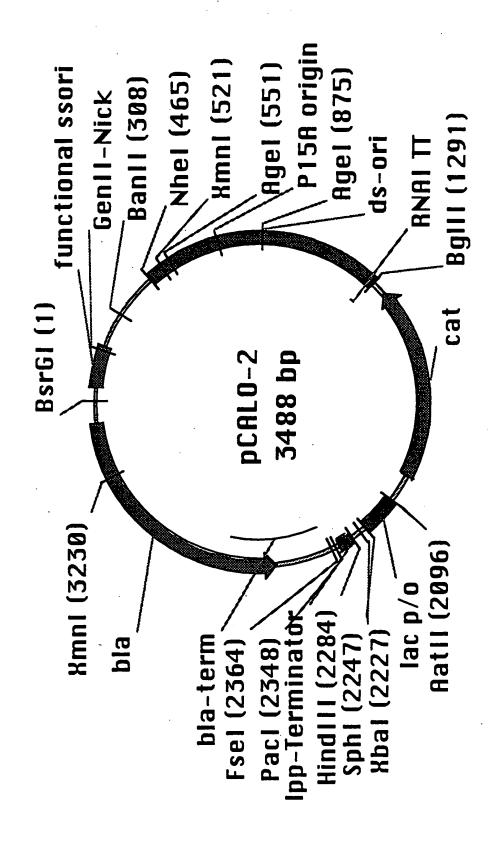


Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

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CCGTTTTAGG GGCAAAATCC CCGGCTTTAG GGCCGAAATC TTAACCAATA AATTGGTTAT TCGAGTAAAA AGCTCATTTT AACAATTTAG TTGTTAAATC 51

GACTCCAACG TCAAAGGGCG ACAAGGTCAA TGTTCCAGTT CCAACTCACA GGTTGAGTGT ACCGAGATAG TGGCTCTATC AAAGAACGTG AAAAGAATAG GTCCACTATT TTTTCTTATC TGGAACAAGA CTTATAAATC GAATATTTAG 101 151

TCACCCTAAT AGTGGGATTA ACGAGAACCA TGCTCTTGGT ATGGCCCACT TACCGGGTGA ATAGTCCCGC TATCAGGGCG TTTTGGCAG AAAACCGTC 201

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GAACCCTAAA CTTGGGATTT CACTAAATCG GTGATTTAGC TGCCGTAAAG ACGGCATTTC GGGGTCGAGG CCCCAGCTCC CAAGTTTTTT GTTCAAAAAA 251

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TTGACGGGGA AAGCCGGCGA ACGTGGCGAG GATTTAGAGC GGGAGCCCCC 301

Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

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| 9.00 | GTGACTGAGC               | GATGCGAGCC               | AGCAAGCTGA               | CGCCGCTCGC               | CTTTACCGAA               |
|------|--------------------------|--------------------------|--------------------------|--------------------------|--------------------------|
| 651  | ACGAACGGGG<br>TGCTTGCCCC | CGGAGATTTC<br>GCCTCTAAAG | CTGGAAGATG               | CCAGGAAGAT<br>GGTCCTTCTA | ACTTAACAGG<br>TGAATTGTCC |
| 701  | GAAGTGAGAG<br>CTTCACTCTC | GGCCGCGGCA               | AAGCCGTTTT<br>TTCGGCAAAA | TCCATAGGCT<br>AGGTATCCGA | CCGCCCCCCT               |
| 751  | GACAAGCATC<br>CTGTTCGTAG | ACGAAATCTG<br>TGCTTTAGAC | ACGCTCAAAT<br>TGCGAGTTTA | CAGTGGTGGC<br>GTCACCACCG | GAAACCCGAC<br>CTTTGGGCTG |
| 801  | AGGACTATAA<br>TCCTGATATT | AGATACCAGG<br>TCTATGGTCC | CGTTTCCCCC               | TGGCGGCTCC               | CTCCTGCGCT               |
|      |                          |                          | AgeI                     |                          |                          |
| 851  | CTCCTGTTCC               | TGCCTTTCGG<br>ACGGAAAGCC | TTTACCGGTG<br>AAATGGCCAC | TCATTCCGCT<br>AGTAAGGCGA | GTTATGGCCG               |
| 901  | CGTTTGTCTC<br>GCAAACAGAG | ATTCCACGCC               | TGACACTCAG               | TTCCGGGTAG<br>AAGGCCCATC | GCAGTTCGCT<br>CGTCAAGCGA |
| 951  | CCAAGCTGGA               | CTGTATGCAC<br>GACATACGTG | GAACCCCCCG               | TTCAGTCCGA               | CCGCTGCGCC               |

Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

| ATGCAAAAGC<br>TACGTTTTCG | AGTCTTGAAG<br>TCAGAACTTC | GTGACTGCGC<br>CACTGACGCG | CAGAGAACCT<br>GTCTCTTGGA | GCAAGAGATT               | н     | GATCTAGCAC<br>CTAGATCGTG | 2020202020               |
|--------------------------|--------------------------|--------------------------|--------------------------|--------------------------|-------|--------------------------|--------------------------|
| ATGC                     | AGTC                     | GTGA                     | CAGA                     | GCAA                     | BgllI | A GATCT<br>T CTAGA       | 0000<br>0000             |
| CCGGAAAGAC<br>GGCCTTTCTG | TAGAGGAGTT<br>ATCTCCTCAA | ACAAGTTTTA<br>TGTTCAAAAT | GTTĠGTAGCT<br>CAACCATCGA | CGTTTTCAGA<br>GCAAAAGTCT |       | CATCTTATTA<br>GTAGAATAAT | AAAAAATTA<br>TTTTTTAAT   |
| TGAGTCCAAC<br>ACTCAGGTTG | GTAATTGATT<br>CATTAACTAA | AACTGAAAGG<br>TTGACTTTCC | GGTTCAAAGA<br>CCAAGTTTCT | GCGGTTTTTT<br>CGCCAAAAAA |       | TCAAGAAGAT<br>AGTTCTTCTA | TAACTGCCTT<br>ATTGACGGAA |
| ACTATCGTCT<br>TGATAGCAGA | GCAGCCACTG               | GTTAAGGCTA<br>CAATTCCGAT | CAGTTACCTC<br>GTCAATGGAG | GCCCTGCAAG<br>CGGGACGTTC |       | CAAAACGATC<br>GTTTTGCTAG | AGGGCACCAA               |
| TTATCCGGTA               | ACCACTGGCA<br>TGGTGACCGT | TCATGCGCCG               | TCCTCCAAGC<br>AGGAGGTTCG | ACGAAAAACC<br>TGCTTTTTGG |       | ACGCGCAGAC<br>TGCGCGTCTG | CAGGCGTTTA<br>GTCCGCAAAT |
| 1001                     | 1051                     | 1101                     | 1151                     | 1201                     |       | 1251                     | 1301                     |
|                          |                          | 9                        | SUBSTITUTE               | SHEET (RUL               | E 26) |                          |                          |

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| AL vector modules and pCAL vectors (co |
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| i di di di di di di di di di di di di di | 1351 | 1351 TGCCACTCAT CGCAGTACTG TTGTAATTCA TTA<br>ACGGTGAGTA GCGTCATGAC AACATTAAGT AAT | CGCAGTACTG               | TTGTAATTCA<br>AACATTAAGT | TTAAGCATTC<br>AATTCGTAAG | TGCCGACATG<br>ACGGCTGTAC |
|--|------|---|--------------------------|--------------------------|--------------------------|--------------------------|
| . •                                      | 1401 | GAAGCCATCA<br>CTTCGGTAGT  | CAAACGGCAT<br>GTTTGCCGTA | GATGAACCTG<br>CTACTTGGAC | AATCGCCAGC<br>TTAGCGGTCG | GGCATCAGCA<br>CCGTAGTCGT |
|  | 1451 | CCTTGTCGCC  | TTGCGTATAA               | TATTTGCCCA               | TAGTGAAAAC<br>ATCACTTTTG | GGGGCGAAG                |
| SUBSTIT                                  | 1501 | AAGTTGTCCA<br>TTCAACAGGT  | TATTGGCTAC               | GTTTAAATCA<br>CAAATTTAGT | AAACTGGTGA<br>TTTGACCACT | AACTCACCCA<br>TTGAGTGGGT |
| UTE SHEET                                | 1551 | GGGATTGGCT<br>CCCTAACCGA  | GAGACGAAAA<br>CTCTGCTTTT | ACATATTCTC<br>TGTATAAGAG | AATAAACCCT<br>TTATTTGGGA | TTAGGGAAAT<br>AATCCCTTTA |
| (RULE 26)                                | 1601 | AGGCCAGGTT<br>TCCGGTCCAA  | TTCACCGTAA               | CACGCCACAT               | CTTGCGAATA<br>GAACGCTTAT | TATGTGTAGA<br>ATACACATCT |
|  | 1651 | AACTGCCGGA<br>TTGACGGCCT  | AATCGTCGTG               | GTATTCACTC               | CAGAGCGATG<br>GTCTCGCTAC | AAAACGTTTC<br>TTTTGCAAAG |
|  | 1701 | AGTTTGCTCA<br>TCAAACGAGT  | TGGAAAACGG<br>ACCTTTTGCC | TGTAACAAGG<br>ACATTGTTCC | GTGAACACTA<br>CACTTGTGAT | TCCCATATCA<br>AGGGTATAGT |

| CATC<br>GTAG   | TTTT<br>AAAA             | TTAT<br>AATA             | ATGC<br>TACG             | TTTT<br>AAAA             | GGTA<br>CCAT             | TAAT<br>ATTA                                | TTCC       |
|--|--------------------------|--------------------------|--------------------------|--------------------------|--------------------------|---|------------|
| AGCATTCATC<br>TCGTAAGTAG   | GCTTATTTTT<br>CGAATAAAAA | GTCTGGTTAT<br>CAGACCAATA | TTTACGATGC<br>AAATGCTACG | TCTCCATTTT<br>AGAGGTAAAA | ACGCCCGGTA<br>TGCGGGCCAT | Aatii<br>~~~~~~<br>GACGTCTAAT<br>CTGCAGATTA | TTATGCTTCC |
| nntinued)<br>ACTCCGGGTG<br>TGAGGCCCAC  | TAAAACTTGT<br>ATTTTGAACA | CAGCTGAACG               | CAAAATGTTC<br>GTTTTACAAG | GTGATTTTTT<br>CACTAAAAAA | CTCAAAAAAT<br>GAGTTTTTTA | AACCTCACCC<br>TTGGAGTGGG                    | GCTTTACACT |
| PCAL vector modules and pCAL vectors (continued) PTTCATT GCCATACGGA ACTO AAAGTAA CGGTATGCCT TGAO | AAAGGCCGGA<br>TTTCCGGCCT | CCGTAATATC<br>GGCATTATAG | TGAAATGCCT<br>ACTTTACGGA | GGTATATCCA<br>CCATATAGGT | ATCTCGATAA<br>TAGAGCTATT | TGAAAGTTGG<br>ACTTTCAACC                    | GGCACCCCAG |
|  | GAATGTGAAT<br>CTTACACTTA | TTTAAAAAGG<br>AAATTTTTCC | AGCAACTGAC<br>TCGTTGACTG | TATCAACGGT<br>ATAGTTGCCA | GCTCCTGAAA<br>CGAGGACTTT | TTCATTATGG                                  | TCACTCATTA |
| Figure 35a: Functional maps and sequences of additional 1751 CCAGCTCACC GTC/GAGTGG CAG           | AGGCGGGCAA<br>TCCGCCCGTT | CTTTACGGTC               | AGGTACATTG<br>TCCATGTAAC | CATTGGGATA<br>GTAACCCTAT | AGCTTCCTTA<br>TCGAAGGAAT | GTGATCTTAT                                  | GTGAGTTAGC |
| ia: Functional<br>1751   | 1801                     | 1851                     | 1901                     | 1951                     | 2001                     | 2051  | 2101       |
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| Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued) |                       |
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| Figure 35a: Functional maps and sequences of additional pCAL vecto                                       | E                     |
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|   | CAC          | CACTCAATCG           | AGTGAGTAAT CCGTGGGGTC CGAAATGTGA AATACGAAGG | CCGTGGGGTC | CGAAATGTGA               | AATACGAAGG               |  |
|---|--------------|----------------------|---|------------|--------------------------|--------------------------|--|
| 2151 GGCTCGTATG TTGTGGAA TTGTGAGCGG ATAACAATTT CACACAGGAA<br>CCGAGCATAC AACACCTT AACACTCGCC TATTGTTAAA GTGTGTCCTT | $\leftarrow$ | CTCGTATG<br>GAGCATAC | •   | TTGTGAGCGG | ATAACAATTT<br>TATTGTTAAA | CACACAGGAA<br>GTGTGTCCTT |  |
| Xbal  | ,            | ÷                    |   | XbaI       |                          | Sphi                     |  |

|      |            | ,          | 2 | }                                      | ?<br>?<br>?<br>? |
|------|------------|------------|---|--|------------------|
| 2201 | ACAGCTATGA | CCATGATTAC | GAATTTCTAG                              | GATTAC GAATTICTAG ACCCCCCCC CGCATGCCAT | CGCATGCCAT       |
|      | TGTCGATACT | GGTACTAATG | CTTAAAGATC                              | ACTAATG CTTAAAGATC TGGGGGGGGG          | GCGTACGGTA       |
|      |            |            |   |  |                  |

|         |         | CCTGTGAAGT                                  |
|---------|---------|---|
| 7777777 | ? ? ? ? | AATGTACGCT ATACGAAGTT ATAAGCTTGA CCTGTGAAGT |
|         |         | ATACGAAGTT                                  |
| •       |         | AATGTACGCT                                  |
|         |         | AACTTCGTAT                                  |
|         |         | 2251  |

| CCTGTGAAGT                                  | GGACACTTCA            | PacI    |
|---|-----------------------|---------|
| AATGTACGCT ATACGAAGTT ATAAGCTTGA CCTGTGAAGT | TATTCGAACT GGACACTTCA |         |
| ATACGAAGTT                                  | TATGCTTCAA            |         |
| AATGTACGCT                                  | TTACATGCGA TATGCTTCAA |         |
| AACTTCGTAT                                  | TTGAAGCATA            |         |
| 2251  |                       |         |
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| ГААТТАА<br>АТТААТТ   |      | FTTGATC   |
|--|------|---|
| TTTGTCTGCC GTTTAATTAA<br>AAACAGACGG CAAATTAATT                       |      | CTCAAGAAGA TCCTTTGATC                           |
| TTTGT  |      | CTCAA   |
| CGACATTTTT TTTGTCTGCC GTTTAATTAA<br>GCTGTAAAAA AAACAGACGG CAAATTAATT | ٠    | CAAAAAGGAT                                      |
| GAAAAATGGC GCAGATTGTG CGACATTTTT<br>CTTTTTACCG CGTCTAACAC GCTGTAAAAA | eI   | GGC CGGCCATTAT CAAAAAGGAT CTCAAGAAGA TCCTTTGATC |
| GAAAAATGGC<br>CTTTTTACCG   | FseI | 29999999999                                     |
| 2301   |      | 2351  |

Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

| 2401 TTTTCTACGG<br>AAAAGATGCC | 2451 TTTGGTCATG<br>AAACCAGTAC | 2501 AAAAATGAAG<br>TTTTTACTTC | 2551 GACAGTTACC            | H 2601 TATTTCGTTC            | M 2651 ATACGGAGG TATGCCCTCC | 2701 CCCACGCTCA<br>GGGTGCGAGT | 2751 GGGCCGAGCG |
|-------------------------------|-------------------------------|-------------------------------|----------------------------|------------------------------|-----------------------------|-------------------------------|-----------------|
| G GETCTGACGC C CCAGACTGCG     | S AGATTATCAA<br>C TCTAATAGTT  | S TTTTAAATCA<br>C AAAATTTAGT  | CAATGCTTAA<br>G GTTACGAATT | C ATCCATAGTT<br>G TAGGTATCAA | G CCTTACCATC                | A CCGGCTCCAG<br>I GGCCGAGGTC  | S CAGAAGTGGT    |
| TCAGTGGAAC<br>AGTCACCTTG      | AAAGGATCTT<br>TTTCCTAGAA      | ATCTAAAGTA<br>TAGATTTCAT      | TCAGTGAGGC<br>AGTCACTCCG   | GCCTGACTCC<br>CGGACTGAGG     | TGGCCCCAGT                  | ATTTATCAGC<br>TAAATAGTCG      | CCTGCAACTT      |
| GAAAACTCAC<br>CTTTTGAGTG      | CACCTAGATC<br>GTGGATCTAG      | TATATGAGTA<br>ATATACTCAT      | ACCTATCTCA<br>TGGATAGAGT   | CCGTCGTGTA                   | GCTGCAATGA<br>CGACGTTACT    | AATAAACCAG<br>TTATTTGGTC      | TATCCGCCTC      |
| GTTAAGGGAT<br>CAATTCCCTA      | CTTTTAAATT<br>GAAAATTTAA      | AACTTGGTCT<br>TTGAACCAGA      | GCGATCTGTC<br>CGCTAGACAG   | GATAACTACG<br>CTATTGATGC     | TACCGCGAGA                  | CCAGCCGGAA<br>GGTCGGCCTT      | CATCCAGTCT      |

Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

| 2801 | ATTAACTGTT<br>TAATTGACAA | GCCGGGAAGC<br>CGGCCCTTCG | TAGAGTAAGT<br>ATCTCATTCA | AGTTCGCCAG<br>TCAAGCGGTC | TTAATAGTTT<br>AATTATCAAA |
|------|--------------------------|--------------------------|--------------------------|--------------------------|--------------------------|
| 2851 | GCGCAACGTT               | GTTGCCATTG               | CTACAGGCAT               | CGTGGTGTCA<br>GCACCACAGT | CGCTCGTCGT<br>GCGAGCAGCA |
| 2901 | TTGGTATGGC               | TTCATTCAGC               | TCCGGTTCCC               | AACGATCAAG<br>TTGCTAGTTC | GCGAGTTACA<br>CGCTCAATGT |
| 2951 | TGATCCCCCA               | TGTTGTGCAA<br>ACAACACGTT | AAAAGCGGTT<br>TTTTCGCCAA | AGCTCCTTCG<br>TCGAGGAAGC | GTCCTCCGAT               |
| 3001 | CGTTGTCAGA               | AGTAAGTTGG<br>TCATTCAACC | CCGCAGTGTT               | ATCACTCATG<br>TAGTGAGTAC | GTTATGGCAG<br>CAATACCGTC |
| 3051 | CACTGCATAA<br>GTGACGTATT | TTCTCTTACT               | GTCATGCCAT               | CCGTAAGATG               | CTTTTCTGTG<br>GAAAAGACAC |
| 3101 | ACTGGTGAGT               | ACTCAACCAA<br>TGAGTTGGTT | GTCATTCTGA               | GAATAGTGTA<br>CTTATCACAT | TGCGGCGACC               |
| 3151 | GAGTTGCTCT<br>CTCAACGAGA | TGCCCGGCGT               | CAATACGGGA<br>GTTATGCCCT | TAATACCGCG               | CCACATAGCA<br>GGTGTATCGT |

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Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

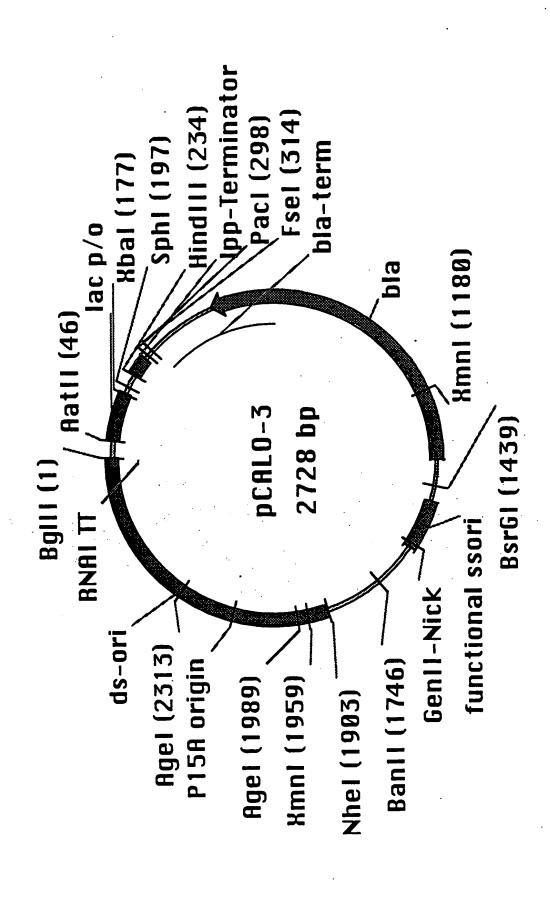
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| •                        |            |            |                       |                          |
|--------------------------|------------|------------|-----------------------|--------------------------|
| GCGAAAACTC<br>CGCTTTTGAG | CCACTCGCGC | TCTGGGTGAG | GGCGACACGG            | GAAGCATTTA<br>CTTCGTAAAT |
| GTTCTTCGGG               | TCGATGTAAC | CACCAGCGTT | AGGGAATAAG            | CAATATTATT               |
|                          | AGCTACATTG | GTGGTCGCAA | TCCCTTATTC            | GTTATAATAA               |
| ATTGGAAAAC               | GAGATCCAGT | CTTTTACTTT | GCCGCAAAAA            | CTTCCTTTTT               |
| TAACCTTTTG               | CTCTAGGTCA | GAAAATGAAA | CGGCGTTTTT            | GAAGGAAAAA               |
| AGTGCTCATC               | TACCGCTGTT | TCCTCAGCAT | AAGGCAAAAT            | AAATGTTGAA TACTCATACT    |
| TCACGAGTAG               | ATGGCGACAA | AGGAGTCGTA | TTCCGTTTTA            | TTTACAACTT ATGAGTATGA    |
| GAACTTTAAA AGTG          | TCAAGGATCT | ACCCAACTGA | CAAAAACAGG AAGGCAAAAT | AAATGTTGAA               |
| CTTGAAATTT TCAC          | AGTTCCTAGA | TGGGTTGACT | GTTTTTGTCC TTCCGTTTTA |                          |
| 3201                     | 3251       | 3301       | 3351                  | 3401                     |
|                          |            | SUBSTITU   | ITE SHEET (           | RULE 28)                 |

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TCAGGGTTAT TGTCTCATGA GCGGATACAT ATTTGAAT AGTCCCAATA ACAGAGTACT CGCCTATGTA TAAACTTA 3451

Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)



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Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

| pCAL | pCALO-3:                   |  |   |                          | †<br>†                                |
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|      | Bglii                      |  |   |                          | AatII                                 |
|      | <b>? ? ? ? ? ? ? ? ? ?</b> |  |   |                          | }<br>}<br>}                           |
| Н    | GATCTCATAA<br>CTAGAGTATT   | CTTCGTATAA<br>GAAGCATATT                       | TGTATGCTAT<br>ACATACGATA                | ACGAAGTTAT<br>TGCTTCAATA | GACGTCTAAT<br>CTGCAGATTA              |
|      |                            |  |   |                          |                                       |
| 21   | GTGAGTTAGC<br>CACTCAATCG   | TCACTCATTA                                     | GGCACCCCAG<br>CCGTGGGGGTC               | GCTTTACACT<br>CGAAATGTGA | TTATGCTTCC<br>AATACGAAGG              |
|      |                            |  |   |                          |                                       |
| 101  | GGCTCGTATG                 | GGCTCGTATG TTGTGTGGAA<br>CCGAGCATAC AACACACCTT | TTGTGAGCGG                              | ATAACAATTT<br>TATTGTTAAA | CACACAGGAA<br>GTGTGTCCTT              |
|      |                            |  |   |                          |                                       |
|      |                            | : .  | XbaI                                    |                          | SphI                                  |
|      |                            | •  | 1 | ?                        | \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ |
| 151  | ACAGCTATGA<br>TGTCGATACT   | CCATGATTAC<br>GGTACTAATG                       | GAATTTCTAG<br>CTTAAAGATC                | ACCCCCCCCC<br>TGGGGGGGGG | CGCATGCCAT<br>GCGTACGGTA              |
|      |                            |  |   |                          |                                       |
|      |                            |  |   | HindIII                  |                                       |
|      |                            |  |   | ? ? ? ? ?                |                                       |
| 201  | AACTTCGTAT                 | AATGTACGCT                                     | ATACGAAGTT                              | ATAAGCTTGA               | CCTGTGAAGT                            |
|      | TTGAAGCATA                 | TTACATGCGA                                     | TATGCTTCAA                              | TATTCGAACT               | GGACACTTCA                            |

Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

|                          |      |                          |                          |                          |                          |                          | -                        |
|--------------------------|------|--------------------------|--------------------------|--------------------------|--------------------------|--------------------------|--------------------------|
| GTTTAATTAA<br>CAAATTAATT |      | TCCTTTGATC               | GTTAAGGGAT<br>CAATTCCCTA | CTTTTAAATT<br>GAAAATTTAA | AACTTGGTCT<br>TTGAACCAGA | GCGATCTGTC<br>CGCTAGACAG | GATAACTACG<br>CTATTGATGC |
| TTTGTCTGCC               |      | CTCAAGAAGA<br>GAGTTCTTCT | GAAACTCAC<br>CTTTTGAGTG  | CACCTAGATC<br>GTGGATCTAG | TATATGAGTA<br>ATATACTCAT | ACCTATCTCA<br>TGGATAGAGT | CCGTCGTGTA               |
| CGACATTTTT<br>GCTGTAAAAA |      | CAAAAAGGAT<br>GTTTTTCCTA | TCAGTGGAAC               | AAAGGATCTT<br>TTTCCTAGAA | ATCTAAAGTA<br>TAGATTTCAT | TCAGTGAGGC               | GCCTGACTCC               |
| GCAGATTGTG<br>CGTCTAACAC | FseI | CGGCCATTAT               | GGTCTGACGC<br>CCAGACTGCG | AGATTATCAA<br>TCTAATAGTT | TTTTAAATCA<br>AAAATTTAGT | CAATGCTTAA<br>GTTACGAATT | ATCCATAGTT<br>TAGGTATCAA |
| GAAAAATGGC<br>CTTTTTACCG | FSET | ပ္ပင္ပ                   | TTTTCTACGG<br>AAAAGATGCC | TTTGGTCATG               | AAAAATGAAG<br>TTTTTACTTC | GACAGTTACC<br>CTGTCAATGG | TATTTCGTTC<br>ATAAAGCAAG |
| 251                      |      | 301                      | 351                      | 401                      | 451                      | 501                      | 551                      |
|                          |      |                          | SUBSTIT                  | UTE SHEET                | (RULE 26)                |                          |                          |

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SUBSTITUTE SHEET (PULE 26)

| CTTTTCTGTG<br>GAAAAGACAC  | TGCGGCGACC               | CCACATAGCA<br>GGTGTATCGT |      | GCGAAAACTC<br>CGCTTTTGAG | CCACTCGCGC<br>GGTGAGCGCG | TCTGGGTGAG<br>AGACCCACTC     | GGCGACACGG<br>CCGCTGTGCC     | GAAGCATTTA   |
|---|--------------------------|--------------------------|------|--------------------------|--------------------------|------------------------------|------------------------------|--------------|
| TAAGATG   | GAATAGTGTA T             | TAATACCGCG CATTATGGCGC   |      | GTTCTTCGGG C             | TCGATGTAAC CAGCTACATTG   | CACCAGCGTT I<br>GTGGTCGCAA A | AGGGAATAAG G<br>TCCCTTATTC C | CAATATTATT G |
| pCAL vector modules and pCAL vectors (continued) PCTTACT GTCATGCCAT CCG' AGAATGA CAGTACGGTA GGC | GTCATTCTGA               | CAATACGGGA<br>GTTATGCCCT | IcmX | ATTGGAAAAC<br>TAACCTTTTG | GAGATCCAGT               | CTTTTACTTT<br>GAAAATGAAA     | GCCGCAAAAA                   | CTTCCTTTTT   |
| Iditional pCAL vector mod<br>TTCTCTTACT<br>AAGAGAATGA   | ACTCAACCAA<br>TGAGTTGGTT | TGCCCGGCGT               |      | AGTGCTCATC<br>TCACGAGTAG | TACCGCTGTT               | TCCTCAGCAT                   | AAGGCAAAAT<br>TTCCGTTTTA     | TACTCATACT   |
| gure 35a: Functional maps and sequences of additional 1001 CACTGCATAA TTCT GTGACGTATT AAGI      | ACTGGTGAGT<br>TGACCACTCA | GAGTTGCTCT<br>CTCAACGAGA |      | GAACTTTAAA<br>CTTGAAATTT | TCAAGGATCT<br>AGTTCCTAGA | ACCCAACTGA<br>TGGGTTGACT     | CAAAAACAGG                   | AAATGTTGAA   |
| gure 35a: Functional<br>1001  | 1051                     | 11.01                    | SUB  | ราบาย เห                 | 1201<br>1301<br>1301     | 1251                         | 1301                         | 1351         |

Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

| CTTCGTAAAT |
|------------|
| GTTATAATAA |
| GAAGGAAAAA |
| ATGAGTATGA |
| TTTACAACTT |

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|      |                          |                          |  | ~        |                          |  |
| 401  | TCAGGGTTAT<br>AGTCCCAATA | TGTCTCATGA<br>ACAGAGTACT | GCGGATACAT<br>CGCCTATGTA                       | TCATGA GCGGATACAT ATTTGAATGT ACATGAAATT        | ACATGAAATT<br>TGTACTTTAA |  |
| .451 | GTAAACGTTA<br>CATTTGCAAT | ATATTTTGTT<br>TATAAAACAA | ATATTTTGTT AAAATTCGCG<br>TATAAAACAA TTTTAAGCGC | TTAAATTTTT GTTAAATCAG<br>AATTTAAAAA CAATTTAGTC | GTTAAATCAG<br>CAATTTAGTC |  |
| 501  | CTCATTTTT                | AACCAATAGG               | CCGAAATCGG                                     | AATAGG CCGAAATCGG CAAAATCCCT TATAAATCAA        | TATAAATCAA               |  |

| TATAAATCAA                                  | CGAGATAGGG TTGAGTGTTG TTCCAGTTTG GAACAAGAGT |
|---|---|
| ATATTTAGTT                                  | GCTCTATCCC AACTCACAAC AAGGTCAAAC CTTGTTCTCA |
| CAAAATCCCT                                  | TTCCAGTTTG<br>AAGGTCAAAC                    |
| AACCAATAGG CCGAAATCGG CAAAATCCCT TATAAATCAA | CGAGATAGGG TTGAGTGTTG TTCCAGTTTG GAACAAGAGT |
| TTGGTTATCC GGCTTTAGCC GTTTTAGGGA ATATTTAGTT | GCTCTATCCC AACTCACAAC AAGGTCAAAC CTTGTTCTCA |
| AACCAATAGG                                  | CGAGATAGGG                                  |
| TTGGTTATCC                                  | GCTCTATCCC                                  |
| CTCAT"I"I"I"I"I                             | AAGAATAGAC                                  |
| GAGTAAAAAA                                  | TTCTTATCTG                                  |
| 1501  | 1551  |
| etiti ite                                   | QUEET (D)                                   |

| ACGTGGA CTCCAACGTC AAAGGGCGAA AAACCGTCTA | GAGGTTGCAG TTTCCCGCTT TTTGGCAGAT |
|--|----------------------------------|
| AAAGGGCGAA                               | GGTTGCAG TTTCCCGCTT I            |
| CTCCAACGTC                               | GAGGTTGCAG                       |
| AGAACGTGGA                               | TCTTGCACCT                       |
| CCACTATTAA                               | GGTGATAATT                       |
| 1601                                     |                                  |

| CCACTAC GAGAACCATC ACCCTAATCA AGTTTTTGG | TCAAAAAACC                              |
|---|---|
| ACCCTAATCA                              | GGTGATG CTCTTGGTAG TGGGATTAGT TCAAAAAAC |
| GAGAACCATC                              | CTCTTGGTAG                              |
| GGCCCACTAC                              | CCGGGTGATG                              |
| TCAGGGCGAT                              | AGTCCCGCTA                              |
| 1651                                    |   |

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| GAGCCCCCGA   | AGGAAGGGAA<br>TCCTTCCCTT     | GCGGTCACGC<br>CGCCAGTGCG     | ACAGGGCGCG<br>TGTCCCGCGC     | TGAGGGTGTC<br>ACTCCCACAG                   |              | GGTGCGTCAG                   | CTGACTCGCT<br>GACTGAGCGA |
|--|------------------------------|------------------------------|------------------------------|--|--------------|------------------------------|--------------------------|
| CTAAAGG<br>GATT VCC  | GTGGCGAGAA A<br>CACCGCTCTT T | GGCAAGTGTA G<br>CCGTTCACAT C | ATGCGCCGCT A<br>TACGCGGCGA T | TTGGCACTGA T                               | AgeI         | AGGCTGCACC G<br>TCCGACGTGG C | TCCTCGCTCA CAGGAGGAGGT G |
| CAL vector modules and pCAL vectors (continued) AAAAGCA CTAAAATCGGA ACC                              | GCCGGCGAAC                   | CTAGGGCGCT                   | GCCGCGCTTA                   | GCTTACTATG<br>CGAATGATAC                   |              | AGGAGAAAAA<br>TCCTCTTTTT     | ATATTCCGCT<br>TATAAGGCGA |
| ditional pCAL vector modu<br>CCGTAAAGCA<br>GGCATTTCGT  | GACGGGGAAA                   | GGAGCGGGCG                   | CACCACACCC<br>GTGGTGTGGG     | GTGTATACTG<br>CACATATGAC                   |              | TTCATGTGGC                   | GATACAGGAT<br>CTATGTCCTA |
| Figure 35a: Functional maps and sequences of additional p<br>1701 GGTCGAGGTG CCGT<br>CCAGCTCCAC GGCA | TTTAGAGCTT<br>AAATCTCGAA     | GAAAGCGAAA<br>CTTTCGCTTT     | TGCGCGTAAC<br>ACGCGCATTG     | NheI<br>~~~~~~<br>TGCTAGCGGA<br>ACGATCGCCT | $\mathbf{H}$ | AGTGAAGTGC                   | CAGAATATGT<br>GTCTTATACA |
| 5a: Functional<br>1701   | 1751                         | 1801                         | 1851                         | 1901                                       |              | 1951                         | 2001                     |
| Figure 3   |                              |                              | SUBSTITU                     | ITE SHEET (RULE 26<br>180 / 204            | )            |                              |                          |

Figure 35a: Functional maps and sequences of additional pCAL vector modules and pCAL vectors (continued)

| 2051 | ACGCTCGGTC<br>TGCGAGCCAG | GTTCGACTGC<br>CAAGCTGACG                   | GGCGAGCGGA<br>CCGCTCGCCT | AATGGCTTAC<br>TTACCGAATG | GAACGGGGGG               |
|------|--------------------------|--|--------------------------|--------------------------|--------------------------|
| 2101 | GAGATTTCCT<br>CTCTAAAGGA | GGAAGATGCC<br>CCTTCTACGG                   | AGGAAGATAC<br>TCCTTCTATG | TTAACAGGGA<br>AATTGTCCCT | AGTGAGAGGG<br>TCACTCTCCC |
| 2151 | CCGCGGCAAA               | GCCGTTTTTC<br>CGGCAAAAAG                   | CATAGGCTCC<br>GTATCCGAGG | GCCCCCCTGA               | CAAGCATCAC<br>GTTCGTAGTG |
| 2201 | GAAATCTGAC<br>CTTTAGACTG | GCTCAAATCA<br>CGAGTTTAGT                   | GTGGTGGCGA               | AACCCGACAG<br>TTGGGCTGTC | GACTATAAAG<br>CTGATATTTC |
| 2251 | ATACCAGGCG<br>TATGGTCCGC | TTTCCCCCTG                                 | GCGGCTCCCT               | CCTGCGCTCT               | CCTGTTCCTG<br>GGACAAGGAC |
| 2301 | CCTTTCGGTT               | AgeI<br>~~~~~~<br>TACCGGTGTC<br>ATGGCCACAG | ATTCCGCTGT               | TATGGCCGCG               | TTTGTCTCAT               |
| 2351 | TCCACGCCTG               | ACACTCAGTT<br>TGTGAGTCAA                   | CCGGGTAGGC               | AGTTCGCTCC<br>TCAAGCGAGG | AAGCTGGACT<br>TTCGACCTGA |

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| AL vector modules and pCAL vec                               |
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| Figure 35a: Functional maps and sequences of additional pCAL |
| map  |
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| AAGTTTTAGT<br>TTCAAAATCA | TGGTAGCTCA GAGAACCTAC<br>ACCATCGAGT CTCTTGGATG | GGTTTTTTCG TTTTCAGAGC AAGAGATTAC |
|--------------------------|--|----------------------------------|
| ပွဲ ပုံ                  |  |                                  |
| CTGAAAGGAC<br>GACTTTCCTG | TTCAAAGAGT<br>AAGTTTCTCA                       | GGTTTTTTCG                       |
| TAAGGCTAAA<br>ATTCCGATTT | GTTACCTCGG<br>CAATGGAGCC                       | CCTGCAAGGC                       |
| 2551                     | 2601   | 2651                             |
|                          |  |                                  |

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AAGAAGATCA TCTTATTA AGAATAAT TTCTTCTAGT AAACGATCTC TTTGCTAGAG 2701

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Figure 35b: List of oligonucleotides used for synthesis of modules

M1: PCR using template

NoVspAatII: TAGACGTC

M2: synthesis

BloxA-A: TATGAGATCTCATAACTTCGTATAATGTACGCTATACG-

**AAGTTAT** 

BloxA-B: TAATAACTTCGTATAGCATACATTATACGAAGTTATG-

**AGATCTCA** 

M3: PCR, NoVspAatII as second oligo

XloxS-muta: CATTTTTGCCCTCGTTATCTACGCATGCGATAACTTCGTA-

TAGCGTACATTATACGAAGTTATTCTAGACATGGTCATAGCTGTTTCCTG

M7-I: PCR

gIIINEW-fow: GGGGGGAATTCGGTGGTGGTGGATCTGCGTGCGCTG-

**AAACGGTTGAAAGTTG** 

gIIINEW-rev: CCCCCCAAGCTTATCAAGACTCCTTATTACG

M7-II: PCR

glllss-fow: GGGGGGGAATTCGGAGGCGGTTCCGGTGGTGGC

M7-III: PCR

glllsupernew-fow: GGGGGGGAATTCGAGCAGAAGCTGATCTCT-

GAGGAGGATCTGTAGGGTGGTGGCTCTGGTTCCGGTGATTTTG

Figure 35b: List of oligonucleotides used for synthesis of modules (continued)

M8: synthesis

lox514-A: CCATAACTTCGTATAATGTACGCTATACGAAGTTATA

lox514-B: AGCTTATAACTTCGTATAGCGTACATTATACGAAGT-

**TATGGCATG** 

M9II: synthesis

M9II-fow: AGCTTGACCTGTGAAGTGAAAAATGGCGCAGATT-

M9II-rev: GTACACCCCCCCAGGCCGGCCCCCCCCCCTTTAA-

TTAAACGGCAGACAAAAAAAAATGTCGCACAATCTGCG

M10II: assembly PCR with template

bla-fow: GGGGGGGTGTACATTCAAATATGTATCCGCTCATG

bla-seq4: GGGTTACATCGAACTGGATCTC

bla1-muta: CCAGTTCGATGTAACCCACTCGCGCACCCAACTGATC-

CTCAGCATCTTTACTTTCACC

blall-muta: ACTCTAGCTTCCCGGCAACAGTTAATAGACTGGATG-

**GAGGCGG** 

bla-NEW: CTGTTGCCGGGAAGCTAGAGTAAG

bla-rev: CCCCCCTTAATTAAGGGGGGGGGCCGGCCATTATCAAA-

**AAGGATCTCAAGAAGATCC** 

M11II/III: PCR, site-directed mutagenesis

Figure 35b: List of oligonucleotides used for synthesis of modules (continued)

f1-fow: GGGGGGGCTAGCACGCCCCTGTAGCGGCGCATTAA

f1-rev: CCCCCCTGTACATGAAATTGTAAACGTTAATATTTTG

f1-t133.muta: GGGCGATGGCCCACTACGAGAACCATCACCCTAATC

#### M12: assembly PCR using template

p15-fow: GGGGGGAGATCTAATAAGATGATCTTCTTGAG

p15-NEWI: GAGTTGGTAGCTCAGAGAACCTACGAAAAACCGCCCTG-

**CAAGGCG** 

p15-NEWII: GTAGGTTCTCTGAGCTACCAACTC

p15-NEWIII: GTTTCCCCCTGGCGGCTCCCTCCTGCGCTCTCCTGTTCCT-

GCC

p15-NEWIV: AGGAGGGAGCCGCCAGGGGAAAC

p15-rev: GACATCAGCGCTAGCGGAGTGTATAC

#### M13: synthesis

BloxXB-A: GATCTCATAACTTCGTATAATGTATGCTATACGAAGTTA-

TTCA

BloxXB-B: GATCTGAATAACTTCGTATAGCATACATTATACGAAGTTA-

**TGAGA** 

M14-Ext2: PCR, site-directed mutagenesis

ColEXT2-fow: GGGGGGGAGATCTGACCAAAATCCCTTAACGTGAG

Col-mutal: GGTATCTGCGCTCTGCTGTAGCCAGTTACCTTCGG

Figure 35b: List of oligonucleotides used for synthesis of modules (continued)

Col-rev: CCCCCCGCTAGCCATGTGAGCAAAAGGCCAGCAA

M17: assembly PCR using template

CAT-1: GGGACGTCGGGTGAGGTTCCAAC

CAT-2: CCATACGGAACTCCGGGTGAGCATTCATC

CAT-3: CCGGAGTTCCGTATGG

CAT-4: ACGTTTAAATCAAAACTGG

CAT-5: CCAGTTTTGATTTAAACGTAGCCAATATGGACAACTTCTTC-

GCCCCGTTTCACTATGGGCAAATATT

CAT-6: GGAAGATCTAGCACCAGGCGTTTAAG

M41: assembly PCR using template

LAC1: GAGGCCGGCCATCGAATGGCGCAAAAC

LAC2: CGCGTACCGTCCTCATGGGAGAAAATAATAC

LAC3: CCATGAGGACGGTACGCGACTGGGCGTGGAGCATCTGGTCGCA-

TTGGGTCACCAGCAAATCCGCTGTTAGCTGGCCCATTAAG

LAC4: GTCAGCGGCGGGATATAACATGAGCTGTCCTCGGTATCGTCG

LAC5: GTTATATCCCGCCGCTGACCACCATCAAAC

LAC6: CATCAGTGAATCGGCCAACGCGCGGGGAGAGGCGGTTTGCGT4TTG-

GGAGCCAGGGTGGTTTTC

LAC7: GGTTAATTAACCTCACTGCCCGCTTTCCAGTCGGGAAACCTGTCGTGCC-

**AGCTGCATCAGTGAATCGGCCAAC** 

M41-MCS-fow: CTAGACTAGTGTTTAAACCGGACCGGGGGGGGGGCTT-

AAGGGGGGGGGG

Figure 35b: List of oligonucleotides used for synthesis of modules (continued)

M41-MCS-rev: CTAGCCCCCCCCCCCCTTAAGCCCCCCCCGGTCCGGT-

TTAAACACTAGT

M41-fow: CTAGACTAGTGTTTAAACCGGACCGGGGGGGGGGCTTAA-

GGGGGGGGGGG

M41-rev: CCCCCCTTAAGTGGGCTGCAAAACAAAACGGCCTCC-

TGTCAGGAAGCCGCTTTTATCGGGTAGCCTCACTGCCCGCTTTCC

M41-A2: GTTGTTGTGCCACGCGGTTAGGAATGTAATTCAGCTCCGC

M41-B1: AACCGCGTGGCACAACAAC

M41-B2: CTTCGTTCTACCATCGACACGACCACGCTGGCACCCAGTTG

M41-C1: GTGTCGATGGTAGAACGAAG

M41-CII: CCACAGCAATAGCATCCTGGTCATCCAGCGGATAGTT-

AATAATCAGCCCACTGACACGTTGCGCGAG

M41-DI: GACCAGGATGCTATTGCTGTGG

M41-DII: CAGCGCGATTTGCTGGTGGCCCAATGCGACCAGATGC

M41-EI: CACCAGCAAATCGCGCTG

M41-EII: CCCGGACTCGGTAATGGCACGCATTGCGCCCAGCGCC

M41-FI: GCCATTACCGAGTCCGGG

M42: synthesis

Eco-H5-Hind-fow: AATTCCACCATCACCATTGACGTCTA

Eco-H5-Hind-rev: AGCTTAGACGTCAATGGTGATGATGGTGG

Figure 36: functional map and sequence of ß-lactamase-MCS module

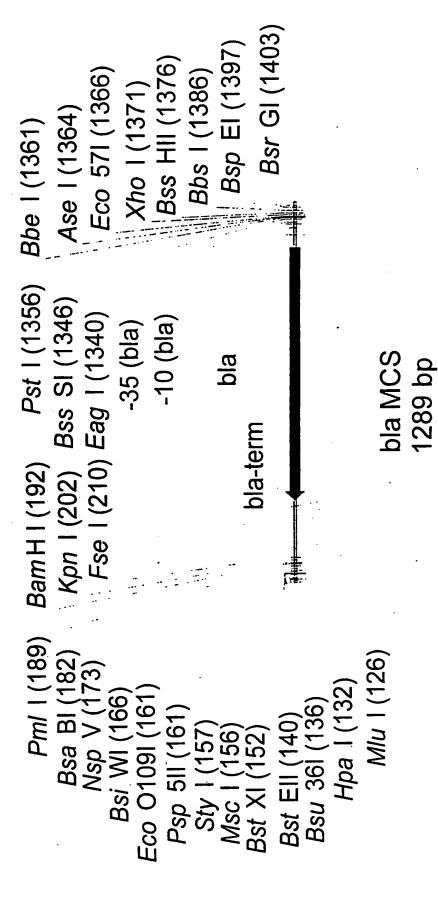


Figure 36: functional map and sequence of B-lactamase-MCS module (continued)

| ٠   |                              |                          |                                  | StyI                           |                              |
|-----|------------------------------|--------------------------|----------------------------------|--------------------------------|------------------------------|
|     |                              |                          |                                  | Psp5II                         |                              |
|     |                              | Bsu36I                   | BstXI                            | Eco01091                       |                              |
|     | HpaI                         | BSTEI                    | Msc                              |                                | BsiwI NspV                   |
| 126 | CGCGTTAACC                   | TCAGGTGACC               | AAGCCCCTGG CCA<br>TTCGGGGACC GGT | AGGTCC                         | C GTACGTTCGA<br>G CATGCAAGCT |
|     |                              | Pml I                    |                                  |                                |                              |
|     | NspVBsaBI                    |                          | KpnI                             | FseI                           |                              |
| 176 | AGATTACCAT C<br>TCTAATGGTA G | CACGTGGATC               | CGGTACCA                         | GG CCGGCCATTA<br>CC GGCCGGTAAT | TCAAAAAGGA<br>AGTTTTTCCT     |
| 226 | TCTCAAGAAG<br>AGAGTTĊTTC     | ATCCTTTGAT<br>TAGGAAACTA | CTTTTCTACG                       | GGGTCTGACG<br>CCCAGACTGC       | CTCAGTGGAA<br>GAGTCACCTT     |
| 276 | CGAAAACTCA<br>GCTTTTGAGT     | CGTTAAGGGA               | TTTTGGTCAT<br>AAAACCAGTA         | GAGATTATCA<br>CTCTAATAGT       | AAAAGGATCT<br>TTTTCCTAGA     |

Figure 36: functional map and sequence of B-lactamase-MCS module (continued)

| AATCTAAAGT<br>TTAGATTTCA | TCAGTGAGGC<br>AGTCACTCCG | GCCTGACTCC<br>CGGACTGAGG | TGGCCCCCAGT | ATTTATCAGC<br>TAAATAGTCG | CCTGCAACTT               | TAGAGTAAGT<br>ATCTCATTCA | CTACAGGCAT<br>GATGTCCGTA |
|--------------------------|--------------------------|--------------------------|-------------|--------------------------|--------------------------|--------------------------|--------------------------|
| GTTTTAAATC               | CAATGCTTAA               | ATCCATAGTT               | GCTTACCATC  | CCGGCTCCAG               | CAGAAGTGGT               | GCCGGGAAGC               | GTTGCCATTG               |
| CAAAATTTAG               | GTTACGAATT               | TAGGTATCAA               | CGAATGGTAG  |                          | GTCTTCACCA               | CGGCCCTTCG               | CAACGGTAAC               |
| TAAAAATGAA               | TGACAGTTAC               | TATTTCGTTC               | ATACGGGAGG  | CCCACGCTCA               | GGGCCGAGCG               | ATTAACTGTT               | GCGCAACGTT               |
| ATTTTTACTT               | ACTGTCAATG               | ATAAAGCAAG               | TATGCCCTCC  |                          | CCCGGCTCGC               | TAATTGACAA               | CGCGTTGCAA               |
| CCTTTTAAAT               | AAACTTGGTC               | GCGATCTGTC               | GATAACTACG  | TACCGCGAGA               | CCAGCCGGAA               | CATCCAGTCT               | TTAATAGTTT               |
| GGAAAATTTA               | TTTGAACCAG               | CGCTAGACAG               | CTATTGATGC  |                          | GGTCGGCCTT               | GTAGGTCAGA               | AATTATCAAA               |
| TCACCTAGAT               | ATATATGAGT<br>TATATACTCA | ACCTATCTCA<br>TGGATAGAGT | CCGTCGTGTA  | GCTGCAATGA<br>CGACGTTACT | AATAAACCAG<br>TTATTTGGTC | TATCCGCCTC               | AGTTCGCCAG<br>TCAAGCGGTC |
| 326                      | 376                      | 426                      | 476         | 526                      | 576                      | 929                      | 919                      |

Figure 36: functional map and sequence of B-lactamase-MCS module (continued)

| TCCGGTTCCC               | AAAAGCGGTT<br>TTTTCGCCAA | CCGCAGTGTT<br>GGCGTCACAA | GTCATGCCAT | GTCATTCTGA<br>CAGTAAGACT | CAATACGGGA<br>GTTATGCCCT | ATTGGAAAAC<br>TAACCTTTTG | GAGATCCAGT<br>CTCTAGGTCA |
|--------------------------|--------------------------|--------------------------|------------|--------------------------|--------------------------|--------------------------|--------------------------|
| TTCATTCAGC               | TGTTGTGCAA               | AGTAAGTTGG               | TTCTCTTACT | ACTCAACCAA               | TGCCCGGCGT               | AGTGCTCATC               | TACCGCTGTT               |
| AAGTAAGTCG               | ACAACACGTT               | TCATTCAACC               | AAGAGAATGA | TGAGTTGGTT               |                          | TCACGAGTAG               | ATGGCGACAA               |
| TTGGTATGGC               | TGATCCCCCA               | CGTTGTCAGA               | CACTGCATAA | ACTGGTGAGT               | GAGTTGCTCT               | GAACTTTAAA               | TCAAGGATCT               |
| AACCATACCG               | ACTAGGGGGT               |                          | GTGACGTATT | TGACCACTCA               | CTCAACGAGA               | CTTGAAATTT               | AGTTCCTAGA               |
| CGCTCGTCGT<br>GCGAGCAGCA | GCGAGTTACA<br>CGCTCAATGT | GTCCTCCGAT               | GTTATGGCAG | CTTTTCTGTG               | TGCGGCGACC               | CCACATAGCA<br>GGTGTATCGT | GCGAAAACTC<br>CGCTTTTGAG |
| CGTGGTGTCA               | AACGATCAAG               | AGCTCCTTCG               | ATCACTCATG | CCGTAAGATG               | GAATAGTGTA               | TAATACCGCG               | GTTCTTCGGG               |
| GCACCACAGT               | TTGCTAGTTC               | TCGAGGAAGC               | TAGTGAGTAC | GGCATTCTAC               | CTTATCACAT               |                          | CAAGAAGCCC               |
| 726                      | 176                      | 826                      | 876        | 926                      | 916                      | 1026                     | 1076                     |

Figure 36: functional map and sequence of ß-lactamase-MCS module (continued)

| CTTTTACTTT<br>GAAAATGAAA          | GCCGCAAAAA<br>CGGCGTTTTT     | CTTCCTTTTT<br>GAAGGAAAAA     | GCGGATACAT                   | XhoI    | BSSHII    | ATGGCTCGAG               |             |
|-----------------------------------|------------------------------|------------------------------|------------------------------|---------|-----------|--------------------------|-------------|
| TCTTCAGCAT CAGAAGTCGTA GECO57I    | AAGGCAAAAT G<br>TTCCGTTTTA C | TACTCATACT C<br>ATGAGTATGA G | TGTCTCATGA G<br>ACAGAGTACT C |         | Bbel Asel | CATTA                    | <b>₩</b> ?  |
| ACCCAACTGA<br>TGGGTTGACT          | CAAAAACAGG<br>GTTTTTGTCC     | AAATGTTGAA<br>TTTACAACTT     | TCAGGGTTAT<br>AGTCCCAATA     | PstI    | BssSI     | ACGAGCTGCA<br>TGCTCGACGT | BspEI BsrGI |
| CCACTCGTGC<br>GGTGAGCACG<br>BSSSI | TCTGGGTGAG                   | GGCGACACGG                   | GAAGCATTTA<br>CTTCGTAAAT     | ·       | EagI      |                          |             |
| TCGATGTAAC<br>AGCTACATTG          | CACCAGCGTT<br>GTGGTCGCAA     | AGGGAATAAG<br>TCCCTTATTC     | СААТАТТАТТ<br>GTTАТААТАА     |         |           | ATTTGAATGT<br>TAAACTTACA | BssHII      |
| 1126                              | 1176                         | 1226                         | 1276                         |         |           | 1326                     | ·           |
|                                   |                              | SUBSTITU                     | JTE SHEET (                  | RULE 26 | 5)        |                          |             |

CATGAAATT GTACTTTAA TCCGGATGTA Figure 36: functional map and sequence of ß-lactamase-MCS module (continued) BbsI CGCTTTGTCT GCGAAACAGA CGCGCTTCAG GCGCGAAGTC Eco57I 1376

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Figure 37: Oligo and primer design for Vκ CDR3 libraries

Figure 37: Oligo and primer design for Vκ CDR3 libraries

G

-3'

F A T/V Y Y C Q
T T T G C G A C T T A T T A T T G C C A

V G V Y Y C
G T G G G C G T G T A T T A T T G C C A

V A V Y Y C

T G G C G G T G T A T T A T T G C

Α C D E G H C A K M N P CAG Q R S T ٧ W Y 80% Q

CA

G

Figure 37: Oligo and primer design for  $V\kappa$  CDR3 libraries

G A C C T

G A C C T

T

|     |    |   |   |   |   |   |            |   |          |         |   |             |       |   |          | •       |          |
|-----|----|---|---|---|---|---|------------|---|----------|---------|---|-------------|-------|---|----------|---------|----------|
| G   | C  | T |   | ******                                  |   |   |            |   | G        | С       | T |             |       |   | G        | C       | Τ        |
|     |    |   |   |   |   |   |            |   |          |         |   |             |       |   |          |         |          |
| G   | Α  | Τ | G | Α                                       | T | G | Α          | T                                       | G        | Α       | T |             |       |   | G        | Α       | T        |
| G   | A  | G |   |   |   |   |            |   | G        | Α       | G |             |       |   | G        | Α       | G        |
| T   | T  | Τ |   | ,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,, |   |   |            | -                                       | T        | T       | T |             |       |   | T        | T       | Τ        |
| G   | G  | T | G | G                                       | T | G | G          | T                                       | G        | G       | T |             |       |   | _        | G       | T        |
| C   | Α  | T |   | •••••                                   |   |   |            |   | С        | Α       | Τ |             |       | · | С        | Α       | T        |
| Α   | T  | T |   |   |   |   |            |   | Α        |         | T |             |       |   | Α        |         | T        |
| Α   | Α  | G |   |   |   |   |            |   |          | Α       | G |             |       |   |          | Α       | G        |
| C   | T  | T |   |   |   |   |            |   | С        |         | T |             |       |   | С        | T       | T        |
| : ' | T  | _ |   |   |   |   |            | ******                                  |          | T       |   |             |       |   |          | T       | G        |
| Α   | Α  | T | Α | Α                                       | T | Α | Α          | T                                       | •        |         |   | •••         |       |   | Α        |         | T        |
|     |    |   |   |   |   |   |            |   |          | ******* |   |             | C     | T | С        |         | T        |
|     | Α  |   |   |   |   |   |            | , q <del>1 4 4 4 8 8 4</del> 7          |          |         | G | <u>:</u>    |       |   | C        | ******  | G        |
| C   | G  |   |   |   |   |   |            | ,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,, | C        |         | T |             |       |   | <u> </u> | G       | T        |
| T   | C  | T | T | C                                       | T | T | C          | T                                       | :<br>2   |         | T | T           | C     | T | T        | ******* | Ţ        |
| A   | С  | T |   |   |   |   |            |   | <u> </u> | С       | Ţ | <u></u>     | ***** |   | Α        | C       | <u>T</u> |
| G   |    | T |   |   |   |   | ********** |   | G        | Ţ       | Ţ | :<br>:<br>: |       |   | G        | Ţ       | Ţ        |
| T   | G  | G |   |   |   |   |            |   | T        |         | G |             | ••••• |   | <u> </u> | G       | G        |
| T   | A  | T | Τ | Α                                       | T |   |            | ********                                | T        | Α       | T | <u> </u>    |       |   | i        | Α       | T        |
| 5   | 0% | Y |   |   |   |   |            |   |          |         |   | 80          | )%    | P |          |         |          |

Figure 37: Oligo and primer design for  $V\kappa$  CDR3 libraries

Figure 38: Oligo and primer design for VA CDR3 libraries

Figure 38: Oligo and primer design for Vλ CDR3 libraries

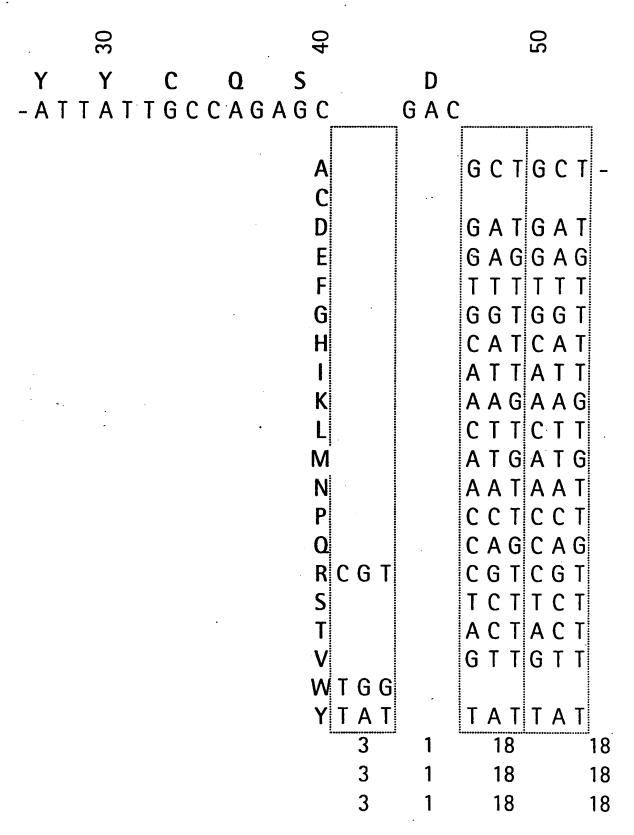


Figure 38: Oligo and primer design for V $\lambda$  CDR3 libraries

| 09                      | 70                     | 80       |
|-------------------------|------------------------|----------|
|                         | 0 0                    | T K L    |
|                         | G G C G G C G G C A    | CGAAGTTA |
| gap gap                 | _                      |          |
| - G C T G C T G C T G C | 1                      |          |
| GATGATGA                | Т                      |          |
| G A G G A G G A G G A   | į                      |          |
| TITITITITI              | Ti '                   |          |
| GGTGGTGGTGG             | T                      |          |
| CATCATCATCA             | T                      |          |
| ATTATTATTAT             | T                      | •        |
| AAGAAGAAGAA             | G                      |          |
| CTTCTTCTTCT.            | T                      |          |
| ATGATGATGAT             | G                      | ·        |
| AATAATAATAA             | <b>T</b> ·             |          |
| CCTCCTCCTCC             | T                      |          |
| C A G C A G C A G C A   | G                      |          |
| CGTCGTCGTCG             | T                      |          |
| TCTTCTTCTTC             | Ţ                      |          |
| ACTACTACTAC             | ]<br>-                 |          |
| GTTGTTGT                |                        |          |
| TATTATTA                |                        |          |
|                         | T Variability 3.32E+05 |          |
| 18 19                   |                        |          |
| 18 18 19<br>18 18 18 19 |                        | •        |
| 18 18 18 19             | 1.UULTUU               |          |

Figure 38: Oligo and primer design for VA CDR3 libraries

Figure 39: functional map of expression vector series pBS13

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Figure 40: Expression data for HuCAL scFvs (pBS13, 30°C)

| % soluble | 7   | Z   | К3         | к4  | ۲٦  | λ2  | 73  |
|-----------|-----|-----|------------|-----|-----|-----|-----|
| H1A       | 61% | 58% | 52%        | 42% | %06 | 61% | %09 |
| H18       | 39% | 48% | <b>%99</b> | 48% | 47% | 39% | 36% |
| H2        | 47% | 57% | 46%        | 49% | 37% | 36% | 45% |
| H3        | 85% | %29 | 76%        | 61% | 80% | 71% | 83% |
| H4        | %69 | 52% | 51%        | 44% | 45% | 33% | 42% |
| H2        | 49% | 49% | 46%        | %29 | 54% | 46% | 47% |
| H6        | %06 | 58% | 54%        | 47% | 45% | 20% | 51% |

| Total amount     |          | Cor  | Ž    | 2        | 7.1  | 72   | 73   |
|------------------|----------|------|------|----------|------|------|------|
| compared to H3K2 | <u>-</u> | لإ   | 2    | <b>†</b> | 1    | 7/5  | 3    |
| H1A              | 289%     | 94%  | 166% | 272%     | 20%  | _    | 78%  |
| H1B              | 219%     | 122% | 89%  |          | 117% | 158% | 101% |
| H2               | 186%     | 223% | 208% |          | 126% | %09  | %26  |
| H3               | 20%      | •    | 71%  | 54%      | 23%  | 130% |      |
| H4               | 37%      | 22%  | %09  | 17%      | 195% | 107% |      |
| H5               | %86      | 201% | 167% | 83%      | 93%  | 128% | 115% |
| 9H               | 65%      | 117% | 89%  | 109%     | 299% | 215% | 278% |

Figure 40: Expression data for HuCAL scFvs (pBS13, 30°C)

| Soluble amount   | <u>,</u> | ć    | Š    |        | 7.1  | 32   | 73       |
|------------------|----------|------|------|--------|------|------|----------|
| compared to H3K2 | <u>-</u> | 2    | 2    | †<br>2 | -₹   | 7.7  | <u>ک</u> |
| H1A              | 191%     | 880% | 121% | 122%   | 26%  | 211% | 76%      |
| H1B              | 124%     | 95%  | 83%  | 107%   | 79%  | 142% | 29%      |
| H2               | 126%     | 204% | 139% | 130%   | %99  | 20%  | 0,002    |
| H3               | 63%      | ı    | 81%  | 49%    | %69  | 143% |          |
| H4               | 40%      | 47%  |      | 54%    | 95%  | 55%  |          |
| H5               | %69      | 158% | 116% | 80%    | 72%  | 84%  | 84%      |
| 9H               | 85%      | 122% | 87%  | 17%    | 162% | 162% | 212%     |
|                  | McPC     |      |      |        |      |      |          |
| soluble          | 38%      |      |      |        |      |      |          |
| %H3k2 total      | 117%     |      |      |        |      |      |          |
| %H3k2 soluble    | %69      | ·    |      |        |      |      |          |

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Hornig, H

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